

Trying 3106016892...Open

```
Welcome to STN International! Enter x:x  
LOGINID:sssptau125rxt  
PASSWORD:  
TERMINAL (ENTER 1, 2, 3, OR ?):2
```

\* \* \* \* \* \* \* \* \* \* \* \* \* \* \* Welcome to STN International \* \* \* \* \* \* \* \* \* \* \*

NEWS	1	Web Page URLs for STN Seminar Schedule - N. America
NEWS	2	Sep 29 The Philippines Inventory of Chemicals and Chemical Substances (PICCS) has been added to CHEMLIST
NEWS	3	Oct 27 New Extraction Code PAX now available in Derwent Files
NEWS	4	Oct 27 SET ABBREVIATIONS and SET PLURALS extended in Derwent World Patents Index files
NEWS	5	Oct 27 Patent Assignee Code Dictionary now available in Derwent Patent Files
NEWS	6	Oct 27 Plasdoc Key Serials Dictionary and Echoing added to Derwent Subscriber Files WPIDS and WPIX
NEWS	7	Nov 29 Derwent announces further increase in updates for DWPII
NEWS	8	Dec 5 French Multi-Disciplinary Database PASCAL Now on STN
NEWS	9	Dec 5 Trademarks on STN - New DEMAS and EUMAS Files
NEWS	10	Dec 15 2001 STN Pricing
NEWS	11	Dec 17 Merged CEABA-VTB for chemical engineering and biotechnology
NEWS	12	Dec 17 Corrosion Abstracts on STN
NEWS	13	Dec 17 SYNTHLINE from Prous Science now available on STN
NEWS	14	Dec 17 The CA Lexicon available in the CAPLUS and CA files
NEWS	15	Jan 05 AIDSILINE is being removed from STN
NEWS	16	Feb 06 Engineering Information Encompass files have new names
NEWS	17	Feb 16 TOXLINE no longer being updated
NEWS EXPRESS		FREE UPGRADE 5.0e FOR STN EXPRESS 5.0 WITH DISCOVER! (WINDOWS) NOW AVAILABLE
NEWS HOURS		STN Operating Hours Plus Help Desk Availability
NEWS INTER		General Internet Information
NEWS LOGIN		Welcome Banner and News Items
NEWS PHONE		Direct Dial and Telecommunication Network Access to STN
NEWS WWW		CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001

=> file reg

COST IN U.S. DOLLARS  
FULL ESTIMATED COST

SINCE FILE ENTRY 0.15  
TOTAL SESSION 0.15

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.  
COPYRIGHT (C) 2001 American Chemical Society (ACS)

STRUCTURE FILE UPDATES: 11 APR 2001 HIGHEST RN 330935-94-9  
DICTIONARY FILE UPDATES: 11 APR 2001 HIGHEST RN 330935-94-9

TSCA INFORMATION NOW CURRENT THROUGH July 8, 2000

Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

Structure search limits have been increased. See HELP SLIMIT  
for details.

=> s neuroserpin

L1 11 NEUROSERPIN

=> d 11 11

L1 ANSWER 11 OF 11 REGISTRY COPYRIGHT 2001 ACS  
RN 179006-00-9 REGISTRY  
CN Axonin 2 (chicken precursor) (9CI) (CA INDEX NAME)  
OTHER NAMES:  
CN 25: PN: WO0053793 FIG: 2 unclaimed sequence  
CN GenBank Z71930-derived protein GI 1359668  
CN Neuroserpin, pre- (chicken)  
FS PROTEIN SEQUENCE  
MF Unspecified  
CI MAN  
SR CA  
LC STN Files: CA, CAPLUS

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*  
\*\*\* USE 'SQD' OR 'SQIDE' FORMATS TO DISPLAY SEQUENCE \*\*\*  
2 REFERENCES IN FILE CA (1967 TO DATE)  
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)

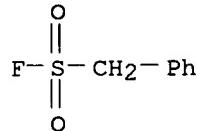
=> s pmsf

L2 1 PMSF

=> d 12

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2001 ACS  
RN 329-98-6 REGISTRY  
CN Benzenemethanesulfonyl fluoride (9CI) (CA INDEX NAME)  
OTHER CA INDEX NAMES:  
CN .alpha.-Toluenesulfonyl fluoride (7CI, 8CI)  
OTHER NAMES:  
CN Phenylmethanesulfonyl fluoride  
CN Phenylmethylsulfonyl fluoride

CN **PMSF**  
 FS 3D CONCORD  
 MF C7 H7 F O2 S  
 CI COM  
 LC STN Files: AGRICOLA, BEILSTEIN\*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA,  
     CANCERLIT, CAOLD, CAPLUS, CASREACT, CHEMCATS, CHEMINFORMRX, CHEMLIST,  
     CSCHEM, DDFU, DRUGU, EMBASE, HODOC\*, IFICDB, IFIPAT, IFIUDB, MEDLINE,  
     MSDS-OHS, NIOSHTIC, PIRA, RTECS\*, SPECINFO, TOXLINE, TOXLIT, USPATFULL  
     (\*File contains numerically searchable property data)  
 Other Sources: DSL\*\*, EINECS\*\*, TSCA\*\*  
     (\*\*Enter CHEMLIST File for up-to-date regulatory information)



726 REFERENCES IN FILE CA (1967 TO DATE)  
 7 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
 727 REFERENCES IN FILE CAPLUS (1967 TO DATE)  
 11 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> s apmsf

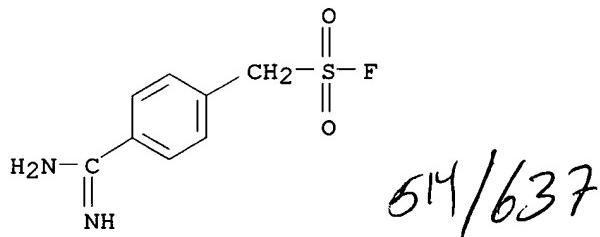
L3           1 APMSF

=> s 13

L4           1 APMSF

=> d 13

L3 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2001 ACS  
 RN 71933-13-6 REGISTRY  
 CN Benzenemethanesulfonyl fluoride, 4-(aminoiminomethyl)- (9CI) (CA INDEX  
     NAME)  
 OTHER NAMES:  
 CN (p-Amidinophenyl)methylsulfonyl fluoride  
 CN **APMSF**  
 CN p-Amidinophenylmethanesulfonyl fluoride  
 FS 3D CONCORD  
 MF C8 H9 F N2 O2 S  
 CI COM  
 LC STN Files: BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CSCHEM, EMBASE,  
     MEDLINE, TOXLIT, USPATFULL



24 REFERENCES IN FILE CA (1967 TO DATE)

=> s antipain

L5 5 ANTIPAIN

=> d 15 5

L5 ANSWER 5 OF 5 REGISTRY COPYRIGHT 2001 ACS

RN 37682-71-6 REGISTRY

CN L-Valinamide,

N2-[(1-carboxy-2-phenylethyl)amino]carbonyl]-L-arginyl-N-[4-[(aminoiminomethyl)amino]-1-formylbutyl]-, compd. with 2,4,6-trinitrophenol (1:2) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN **Antipain dipicrate**

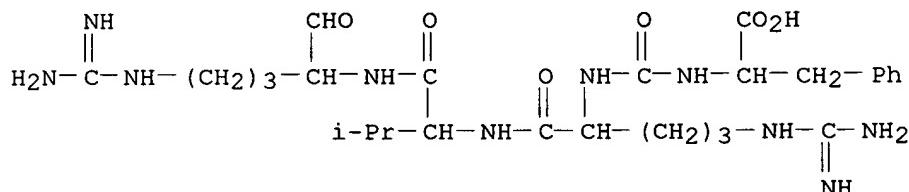
MF C27 H44 N10 O6 . 2 C6 H3 N3 O7

LC STN Files: CA, CAPLUS

CM 1

CRN 37691-11-5

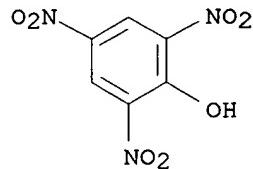
CMF C27 H44 N10 O6



CM 2

CRN 88-89-1

CMF C6 H3 N3 O7



1 REFERENCES IN FILE CA (1967 TO DATE)

1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

=> s antithrombin

L6 170 ANTITHROMBIN

=> s 16 170

MISSING OPERATOR

=> d 16 170

L6 ANSWER 170 OF 170 REGISTRY COPYRIGHT 2001 ACS  
RN 9000-94-6 REGISTRY  
CN **Antithrombin (9CI)** (CA INDEX NAME)  
OTHER NAMES:  
CN **Antithrombin III**  
CN Heparin cofactor  
CN Heparin cofactor B  
CN Org 10849  
CN Thrombin inhibitor  
AR 90170-80-2  
DR 9041-91-2, 52014-67-2  
MF Unspecified  
CI PMS, COM, MAN  
PCT Manual registration  
LC STN Files: AGRICOLA, AIDSLINE, ANABSTR, BIOBUSINESS, BIOSIS, BIOTECHNO,  
CA, CANCERLIT, CAPLUS, CBNB, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM,  
DDFU,  
DRUGNL, DRUGPAT, DRUGU, DRUGUPDATES, EMBASE, IFICDB, IFIPAT, IFIUDB,  
IPA, MEDLINE, MSDS-OHS, NIOSHTIC, PHAR, PROMT, RTECS\*, TOXLINE, TOXLIT,  
USAN, USPATFULL  
(\*File contains numerically searchable property data)  
Other Sources: EINECS\*\*  
(\*\*Enter CHEMLIST File for up-to-date regulatory information)  
  
\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*  
4265 REFERENCES IN FILE CA (1967 TO DATE)  
471 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
4278 REFERENCES IN FILE CAPLUS (1967 TO DATE)

=> s leupeptin

L7 11 LEUPEPTIN

=> d 17 1

L7 ANSWER 1 OF 11 REGISTRY COPYRIGHT 2001 ACS  
RN 81458-06-2 REGISTRY  
CN **Peptidase, leupeptin (9CI)** (CA INDEX NAME)  
OTHER NAMES:  
CN **Leupeptin peptidase**  
CN **Leupeptin-inactivating enzyme**  
MF Unspecified  
CI MAN  
LC STN Files: BIOSIS, CA, CAPLUS, TOXLIT  
  
\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*  
7 REFERENCES IN FILE CA (1967 TO DATE)  
7 REFERENCES IN FILE CAPLUS (1967 TO DATE)

=> d 178\

'L78\' IS NOT A VALID FORMAT FOR FILE 'REGISTRY'

The following are valid formats:

Substance information can be displayed by requesting individual fields or predefined formats. The predefined substance formats

are: (RN = CAS Registry Number)

REG - RN  
SAM - Index Name, MF, and structure - no RN  
FIDE - All substance data, except sequence data  
IDE - FIDE, but only 50 names  
SQIDE - IDE, plus sequence data  
SQIDE3 - Same as SQIDE, but 3-letter amino acid codes are used  
SQD - Protein sequence data, includes RN  
SQD3 - Same as SQD, but 3-letter amino acid codes are used  
SQN - Protein sequence name information, includes RN

Any CA File format may be combined with any substance format to obtain CA references citing the substance. The substance formats must be cited first. The CA File predefined formats are:

ABS -- Abstract  
APPS -- Application and Priority Information  
BIB -- CA Accession Number, plus Bibliographic Data  
CAN -- CA Accession Number  
CBIB -- CA Accession Number, plus Bibliographic Data (compressed)  
IND -- Index Data  
IPC -- International Patent Classification  
PATS -- PI, SO  
STD -- BIB, IPC, and NCL

IABS --ABS, indented, with text labels  
IBIB -- BIB, indented, with text labels  
ISTD -- STD format, indented

OBIB ----- AN, plus Bibliographic Data (original)  
OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations  
SIBIB ----- IBIB, no citations

The ALL format gives FIDE BIB ABS IND RE, plus sequence data when it is available.

The MAX format is the same as ALL.

The IALL format is the same as ALL with BIB ABS and IND indented, with text labels.

For additional information, please consult the following help messages:

HELP DFIELDS -- To see a complete list of individual display fields.  
HELP FORMATS -- To see detailed descriptions of the predefined formats.  
ENTER DISPLAY FORMAT (IDE):bib

'BIB' IS NOT A VALID FORMAT FOR FILE 'REGISTRY'

The following are valid formats:

Substance information can be displayed by requesting individual fields or predefined formats. The predefined substance formats are: (RN = CAS Registry Number)

REG - RN  
SAM - Index Name, MF, and structure - no RN  
FIDE - All substance data, except sequence data  
IDE - FIDE, but only 50 names  
SQIDE - IDE, plus sequence data  
SQIDE3 - Same as SQIDE, but 3-letter amino acid codes are used

SQD - Protein sequence data, includes RN  
SQD3 - Same as SQD, but 3-letter amino acid codes are used  
SQN - Protein sequence name information, includes RN

Any CA File format may be combined with any substance format to obtain CA references citing the substance. The substance formats must be cited first. The CA File predefined formats are:

ABS -- Abstract  
APPS -- Application and Priority Information  
BIB -- CA Accession Number, plus Bibliographic Data  
CAN -- CA Accession Number  
CBIB -- CA Accession Number, plus Bibliographic Data (compressed)  
IND -- Index Data  
IPC -- International Patent Classification  
PAT5 -- PI, SO  
STD -- BIB, IPC, and NCL

IABS -- ABS, indented, with text labels  
IBIB -- BIB, indented, with text labels  
ISTD -- STD format, indented

OBIB ----- AN, plus Bibliographic Data (original)  
OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations  
SIBIB ----- IBIB, no citations

The ALL format gives FIDE BIB ABS IND RE, plus sequence data when it is available.

The MAX format is the same as ALL.

The IALL format is the same as ALL with BIB ABS and IND indented, with text labels.

For additional information, please consult the following help messages:

HELP DFIELDS -- To see a complete list of individual display fields.  
HELP FORMATS -- To see detailed descriptions of the predefined formats.  
ENTER DISPLAY FORMAT (IDE):abs

'ABS' IS NOT A VALID FORMAT FOR FILE 'REGISTRY'

The following are valid formats:

Substance information can be displayed by requesting individual fields or predefined formats. The predefined substance formats are: (RN = CAS Registry Number)

REG - RN  
SAM - Index Name, MF, and structure - no RN  
FIDE - All substance data, except sequence data  
IDE - FIDE, but only 50 names  
SQIDE - IDE, plus sequence data  
SQIDE3 - Same as SQIDE, but 3-letter amino acid codes are used  
SQD - Protein sequence data, includes RN  
SQD3 - Same as SQD, but 3-letter amino acid codes are used  
SQN - Protein sequence name information, includes RN

Any CA File format may be combined with any substance format to obtain CA references citing the substance. The substance formats must be cited first. The CA File predefined formats are:

ABS -- Abstract  
APPS -- Application and Priority Information  
BIB -- CA Accession Number, plus Bibliographic Data  
CAN -- CA Accession Number  
CBIB -- CA Accession Number, plus Bibliographic Data (compressed)  
IND -- Index Data  
IPC -- International Patent Classification  
PATS -- PI, SO  
STD -- BIB, IPC, and NCL

IABS --ABS, indented, with text labels  
IBIB -- BIB, indented, with text labels  
ISTD -- STD format, indented

OBIB ----- AN, plus Bibliographic Data (original)  
OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations  
SIBIB ----- IBIB, no citations

The ALL format gives FIDE BIB ABS IND RE, plus sequence data when it is available.

The MAX format is the same as ALL.

The IALL format is the same as ALL with BIB ABS and IND indented, with text labels.

For additional information, please consult the following help messages:

HELP DFIELDS -- To see a complete list of individual display fields.  
HELP FORMATS -- To see detailed descriptions of the predefined formats.  
ENTER DISPLAY FORMAT (IDE):

ENTER DISPLAY FORMAT (IDE):

ENTER DISPLAY FORMAT (IDE):ide

L7 ANSWER 1 OF 11 REGISTRY COPYRIGHT 2001 ACS  
RN 81458-06-2 REGISTRY  
CN Peptidase, leupeptin (9CI) (CA INDEX NAME)  
OTHER NAMES:  
CN Leupeptin peptidase  
CN Leupeptin-inactivating enzyme  
MF Unspecified  
CI MAN  
LC STN Files: BIOSIS, CA, CAPLUS, TOXLIT

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*  
7 REFERENCES IN FILE CA (1967 TO DATE)  
7 REFERENCES IN FILE CAPLUS (1967 TO DATE)

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

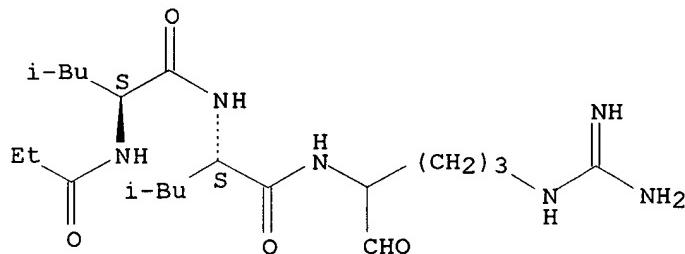
FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001  
L1 11 S NEUROSERPIN  
L2 1 S PMSF  
L3 1 S APMSF  
L4 1 S L3

L5            5 S ANTIPAIN  
L6            170 S ANTITHROMBIN  
L7            11 S LEUPEPTIN

=> d 17 11

L7    ANSWER 11 OF 11    REGISTRY    COPYRIGHT 2001 ACS  
RN    24125-15-3    REGISTRY  
CN    Valeramide, N-(1-formyl-4-guanidinobutyl)-4-methyl-2-(4-methyl-2-propionamidovaleramido)-, monohydrochloride, stereoisomer (8CI)    (CA  
INDEX  
    NAME)  
OTHER CA INDEX NAMES:  
CN    Argininal, N2-[N-(N-propionyl-L-leucyl)-L-leucyl]-, monohydrochloride,  
DL-  
    (8CI)  
OTHER NAMES:  
CN    Leupeptin propyl-LL hydrochloride  
FS    STEREOSEARCH  
DR    36794-11-3  
MF    C21 H40 N6 O4 . Cl H  
LC    STN Files: CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXLIT, USPATFULL  
CRN    (24365-46-6)

Absolute stereochemistry.



● HCl

4 REFERENCES IN FILE CA (1967 TO DATE)  
4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

=> d dichlorocoumarin

'DICHLOROCOUMARIN' IS NOT A VALID FORMAT FOR FILE 'REGISTRY'

The following are valid formats:

Substance information can be displayed by requesting individual fields or predefined formats. The predefined substance formats are: (RN = CAS Registry Number)

REG    - RN  
SAM    - Index Name, MF, and structure - no RN  
FIDE    - All substance data, except sequence data  
IDE    - FIDE, but only 50 names  
SQIDE    - IDE, plus sequence data

SQIDE3 - Same as SQIDE, but 3-letter amino acid codes are used  
SQD - Protein sequence data, includes RN  
SQD3 - Same as SQD, but 3-letter amino acid codes are used  
SQN - Protein sequence name information, includes RN

Any CA File format may be combined with any substance format to obtain CA references citing the substance. The substance formats must be cited first. The CA File predefined formats are:

ABS -- Abstract  
APPS -- Application and Priority Information  
BIB -- CA Accession Number, plus Bibliographic Data  
CAN -- CA Accession Number  
CBIB -- CA Accession Number, plus Bibliographic Data (compressed)  
IND -- Index Data  
IPC -- International Patent Classification  
PATS -- PI, SO  
STD -- BIB, IPC, and NCL

IABS --ABS, indented, with text labels  
IBIB -- BIB, indented, with text labels  
ISTD -- STD format, indented

OBIB ----- AN, plus Bibliographic Data (original)  
OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations  
SIBIB ----- IBIB, no citations

The ALL format gives FIDE BIB ABS IND RE, plus sequence data when it is available.

The MAX format is the same as ALL.

The IALL format is the same as ALL with BIB ABS and IND indented, with text labels.

For additional information, please consult the following help messages:

HELP DFIELDS -- To see a complete list of individual display fields.  
HELP FORMATS -- To see detailed descriptions of the predefined formats.  
ENTER DISPLAY FORMAT (IDE):ide

L7 ANSWER 1 OF 11 REGISTRY COPYRIGHT 2001 ACS  
RN 81458-06-2 REGISTRY  
CN Peptidase, leupeptin (9CI) (CA INDEX NAME)  
OTHER NAMES:  
CN Leupeptin peptidase  
CN Leupeptin-inactivating enzyme  
MF Unspecified  
CI MAN  
LC STN Files: BIOSIS, CA, CAPLUS, TOXLIT

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*  
7 REFERENCES IN FILE CA (1967 TO DATE)  
7 REFERENCES IN FILE CAPLUS (1967 TO DATE)

=> s dichlorocoumarin

L8 9 DICHLOROCOUMARIN

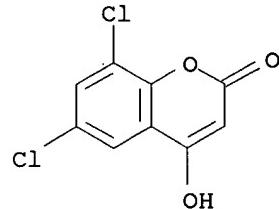
=> d 18 709

9 ANSWERS ARE AVAILABLE. SPECIFIED ANSWER NUMBER EXCEEDS ANSWER SET SIZE

The answer numbers requested are not in the answer set.

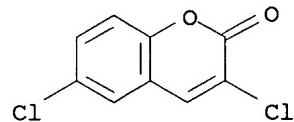
ENTER ANSWER NUMBER OR RANGE (1):7-9

L8 ANSWER 7 OF 9 REGISTRY COPYRIGHT 2001 ACS  
RN 36051-82-8 REGISTRY  
CN 2H-1-Benzopyran-2-one, 6,8-dichloro-4-hydroxy- (9CI) (CA INDEX NAME)  
OTHER CA INDEX NAMES:  
CN Coumarin, 6,8-dichloro-4-hydroxy- (7CI)  
OTHER NAMES:  
CN **4-Hydroxy-6,8-dichlorocoumarin**  
CN 6,8-Dichloro-4-hydroxycoumarin  
FS 3D CONCORD  
MF C9 H4 Cl2 O3  
LC STN Files: BEILSTEIN\*, CA, CAOLD, CAPLUS, CASREACT, CHEMCATS  
(\*File contains numerically searchable property data)



4 REFERENCES IN FILE CA (1967 TO DATE)  
4 REFERENCES IN FILE CAPLUS (1967 TO DATE)  
1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L8 ANSWER 8 OF 9 REGISTRY COPYRIGHT 2001 ACS  
RN 20882-68-2 REGISTRY  
CN 2H-1-Benzopyran-2-one, 3,6-dichloro- (9CI) (CA INDEX NAME)  
OTHER CA INDEX NAMES:  
CN Coumarin, 3,6-dichloro- (7CI, 8CI)  
OTHER NAMES:  
CN **3,6-Dichlorocoumarin**  
FS 3D CONCORD  
MF C9 H4 Cl2 O2  
LC STN Files: BEILSTEIN\*, CA, CAOLD, CAPLUS  
(\*File contains numerically searchable property data)



2 REFERENCES IN FILE CA (1967 TO DATE)  
2 REFERENCES IN FILE CAPLUS (1967 TO DATE)  
1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L8 ANSWER 9 OF 9 REGISTRY COPYRIGHT 2001 ACS  
RN 2199-91-9 REGISTRY  
CN 2H-1-Benzopyran-2-one, 3-acetyl-6,8-dichloro- (9CI) (CA INDEX NAME)  
OTHER CA INDEX NAMES:

CN Coumarin, 3-acetyl-6,8-dichloro- (6CI, 7CI, 8CI)

OTHER NAMES:

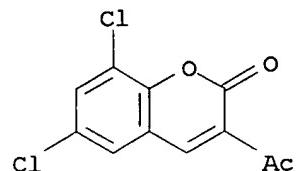
CN 3-Acetyl-6,8-dichlorocoumarin

FS 3D CONCORD

MF C11 H6 Cl2 O3

LC STN Files: BEILSTEIN\*, CA, CAOLD, CAPLUS, CASREACT, CHEMCATS, CSCHEM

(\*File contains numerically searchable property data)



4 REFERENCES IN FILE CA (1967 TO DATE)

4 REFERENCES IN FILE CAPLUS (1967 TO DATE)

2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> s tlck

L9 2 TLCK

=> d 19 1-2

L9 ANSWER 1 OF 2 REGISTRY COPYRIGHT 2001 ACS

RN 23877-38-5 REGISTRY

CN Benzenesulfonamide, N-[(1S)-1-(chloroacetyl)-3-methylbutyl]-4-methyl- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Benzenesulfonamide, N-[1-(chloroacetyl)-3-methylbutyl]-4-methyl-, (S)-

CN p-Toluenesulfonamide, N-[1-(chloroacetyl)-3-methylbutyl]-, stereoisomer (8CI)

OTHER NAMES:

CN L-1-Tosylamide-2-leucyl chloromethyl ketone

CN TLCK

CN Tosyl-L-leucyl chloromethyl ketone

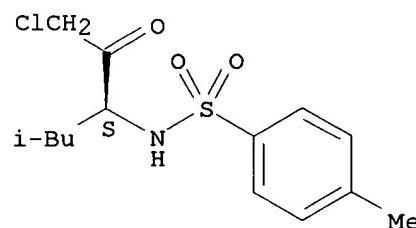
FS STEREOSEARCH

MF C14 H20 Cl N O3 S

LC STN Files: AGRICOLA, BEILSTEIN\*, BIOSIS, CA, CAPLUS, TOXLIT, USPATFULL

(\*File contains numerically searchable property data)

Absolute stereochemistry.



9 REFERENCES IN FILE CA (1967 TO DATE)

9 REFERENCES IN FILE CAPLUS (1967 TO DATE)

L9 ANSWER 2 OF 2 REGISTRY COPYRIGHT 2001 ACS

RN 2364-87-6 REGISTRY

CN Benzenesulfonamide, N-[(1S)-5-amino-1-(chloroacetyl)pentyl]-4-methyl-  
 (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Benzenesulfonamide, N-[5-amino-1-(chloroacetyl)pentyl]-4-methyl-, (S)-

CN p-Toluenesulfonamide, N-[5-amino-1-(chloroacetyl)pentyl]-, L- (8CI)

OTHER NAMES:

CN .alpha.-N-(p-Tosyl)-L-lysyl chloromethyl ketone

CN 1-Chloro-3-tosylamido-7-amino-L-2-heptanone

CN L-1-Chloro-3-tosylamido-7-amino-2-heptanone

CN N-.alpha.-p-Tosyl-L-lysine chloromethylketone

CN N-.alpha.-Tosyl-L-lysyl-chloromethyl ketone

CN N-Tosyl-L-lysine chloromethyl ketone

CN N-Tosyl-L-lysyl chloromethyl ketone

CN N.alpha.-p-Tosyl-L-lysine chloromethyl ketone

CN N.alpha.-p-Tosyl-L-lysylchloromethyl ketone

CN N.alpha.-Tosyl-L-lysine chloromethyl ketone

CN N.alpha.-Tosyl-L-lysyl chloromethyl ketone

CN **TLCK**

CN Tosyl-L-lysine chloromethyl ketone

CN Tosyllysine chloromethyl ketone

CN Tosyllysyl chloromethyl ketone

FS STEREOSEARCH

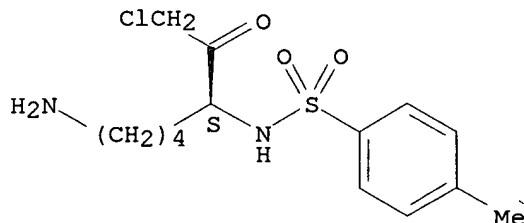
DR 130021-39-5, 3414-37-7

MF C14 H21 Cl N2 O3 S

CI COM

LC STN Files: AGRICOLA, BEILSTEIN\*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA,  
 CAOLD, CAPLUS, CSCHEM, DDFU, DRUGU, EMBASE, NIOSHTIC, RTECS\*, TOXLINE,  
 TOXLIT, USPATFULL  
 (\*File contains numerically searchable property data)

Absolute stereochemistry.



364 REFERENCES IN FILE CA (1967 TO DATE)  
 6 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
 365 REFERENCES IN FILE CAPLUS (1967 TO DATE)  
 1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	57.68	57.83

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(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1	11 S NEUROSERPIN
L2	1 S PMSF
L3	1 S APMSF
L4	1 S L3
L5	5 S ANTIPAIN
L6	170 S ANTITHROMBIN
L7	11 S LEUPEPTIN
L8	9 S DICHLOROCOUMARIN
L9	2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

=> file ca

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.62	58.45

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001  
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FILE COVERS 1967 - 5 Apr 2001 VOL 134 ISS 16  
FILE LAST UPDATED: 5 Apr 2001 (20010405/ED)

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CAS, is strictly prohibited.

=> s 11

L10 17 L1

=> e neuroserpin

E1	1	NEUROSECRETING/BI
E2	1	NEUROSERETORY/BI
E3	17	--> NEUROSERPIN/BI
E4	2	NEUROSERPINS/BI
E5	67	NEUROSES/BI
E6	3	NEUROSHELL/BI
E7	2	NEUROSIGNAL/BI
E8	2	NEUROSIGNALLING/BI
E9	10	NEUROSIN/BI
E10	371	NEUROSIS/BI
E11	1	NEUROSISLIKE/BI
E12	1	NEUROSKELETAL/BI

=> s 12

L11 728 L2

=> s 13

L12 24 L3

=> s 15

L13 343 L5

=> s 16

L14 4326 L6

=> s 17

L15 171 L7

=> s 18

L16 16 L8

=> s 19

L17 372 L9

=> e neuropathy

E1	3	NEUROPATHWAY/BI
E2	6	NEUROPATHWAYS/BI
E3	5490	--> NEUROPATHY/BI
E4	1	NEUROPATHYS/BI
E5	1	NEUROPATTERNING/BI
E6	1	NEUROPATHY/BI
E7	1	NEUROPDPTIDE/BI
E8	2	NEUROPAEDIATRIC/BI
E9	3	NEUROPAEDIATRICS/BI
E10	1	NEUROPEITHELIUM/BI
E11	1	NEUROPELAGIC/BI

E12            3        NEUROOPENIA/BI

=> s e3

L18            5490 NEUROPATHY/BI

=> e epilepsy

E1	3	EPILEPSIS/BI
E2	1	EPILEPSRY/BI
E3	8845 -->	EPILEPSY/BI
E4	1	EPILEPSYE/BI
E5	1	EPILEPTGENIC/BI
E6	1	EPILEPTI/BI
E7	3510	EPILEPTIC/BI
E8	4	EPILEPTICAL/BI
E9	237	EPILEPTICS/BI
E10	696	EPILEPTICUS/BI
E11	1	EPILEPTICY/BI
E12	1392	EPILEPTIFORM/BI

=> s e3-e12

8845	EPILEPSY/BI
1	EPILEPSYE/BI
1	EPILEPTGENIC/BI
1	EPILEPTI/BI
3510	EPILEPTIC/BI
4	EPILEPTICAL/BI
237	EPILEPTICS/BI
696	EPILEPTICUS/BI
1	EPILEPTICY/BI
1392	EPILEPTIFORM/BI
L19	10805 (EPILEPSY/BI OR EPILEPSYE/BI OR EPILEPTGENIC/BI OR EPILEPTI/BI OR EPILEPTIC/BI OR EPILEPTICAL/BI OR EPILEPTICS/BI OR EPILEPTICU S/BI OR EPILEPTICY/BI OR EPILEPTIFORM/BI)

=> e seizure

E1	1	SEIZUROGENIC/BI
E2	3	SEIZURAL/BI
E3	9693 -->	SEIZURE/BI
E4	9	SEIZURED/BI
E5	8	SEIZURELIKE/BI
E6	10034	SEIZURES/BI
E7	2	SEIZURGENIC/BI
E8	11	SEIZURING/BI
E9	4	SEIZUROGENIC/BI
E10	1	SEIZURS/BI
E11	19	SEJ/BI
E12	1	SEJANUS/BI

=> s e3-e6

9693	SEIZURE/BI
9	SEIZURED/BI
8	SEIZURELIKE/BI
10034	SEIZURES/BI
L20	14562 (SEIZURE/BI OR SEIZURED/BI OR SEIZURELIKE/BI OR SEIZURES/BI)

=> e hypoxia

E1           1       HYPOXENOLITHS/BI  
E2           3       HYPOXI/BI  
E3    24801 --> HYPOXIA/BI  
E4           1       HYPOXIAIGNIFICANTLY/BI  
E5           1       HYPOXIAIN/BI  
E6           1       HYPOXIAINDUCED/BI  
E7           1       HYPOXIACHEMIA/BI  
E8           2       HYPOXIAL/BI  
E9           1       HYPOXIAM/BI  
E10          1       HYPOXIANORMOXIA/BI  
E11          1       HYPOXIAPNITRIC/BI  
E12          17      HYPOXIAS/BI

=> s e3

L21          24801 HYPOXIA/BI

=> e stroke

E1           3       STROKAN/BI  
E2           1       STROKAR/BI  
E3    11880 --> STROKE/BI  
E4           1       STROKE1/BI  
E5           1       STROKECYCLE/BI  
E6           31      STROKED/BI  
E7           16      STROKELIKE/BI  
E8           6       STROKEPRONE/BI  
E9           3       STROKER/BI  
E10          895     STROKES/BI  
E11          1       STROKESTOWN/BI  
E12          1       STROKILACEUM/BI

=> s e3

L22          11880 STROKE/BI

=> s 117 and 118

L23          0 L17 AND L18

=> s 117 and 119

L24          0 L17 AND L19

=> s 117 and 120

L25          0 L17 AND L20

=> s 117 and 121

L26          0 L17 AND L21

=> s 117 and 122

L27          0 L17 AND L22

=> s 116 and 118

L28          0 L16 AND L18

=> s 118 or 119 or 120

L29          25960 L18 OR L19 OR L20

=> s 129 and l16

L30 O L29 AND L16

=> s 129 and l15

L31 O L29 AND L15

=> s 129 and l14

L32 4 L29 AND L14

=> d 132 1-4

L32 ANSWER 1 OF 4 CA COPYRIGHT 2001 ACS

AN 133:320515 CA

TI Clinical and biochemical characteristics of congenital disorder of glycosylation type Ic, the first recognized endoplasmic reticulum defect in N-glycan synthesis

AU Grunewald, S.; Imbach, T.; Huijben, K.; Rubio-Gozalbo, M. E.; Verrrips, A.; De Klerk, J. B. C.; Stroink, H.; De Rijk-Van Andel, J. F.; Van Hove, J.

L. K.; Wendel, U.; Matthijs, G.; Hennet, T.; Jaeken, J.; Wevers, R. A.

CS Department of Pediatrics, Heinrich-Heine University Dusseldorf, Dusseldorf, Germany

SO Ann. Neurol. (2000), 47(6), 776-781

CODEN: ANNED3; ISSN: 0364-5134

PB Lippincott Williams & Wilkins

DT Journal

LA English

RE.CNT 29

RE

(3) Burda, P; J Clin Invest 1998, V102, P647 CA

(5) de Koning, T; Biochem Biophys Res Commun 1998, V245, P38 CA

(7) Grunewald, S; Biochem Biophys Acta 1999, V1455, P54 CA

(9) Imbach, T; Proc Natl Acad Sci USA 1999, V96, P6982 CA

(11) Jaeken, J; Am J Hum Genet 1998, V62, P1535 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L32 ANSWER 2 OF 4 CA COPYRIGHT 2001 ACS

AN 133:15850 CA

TI Cerebral venous sinus thrombosis associated with hepatic cirrhosis

AU Singhal, A. B.; Buonanno, F.; Rordorf, G.

CS Department of Neurology, VBK 802, Massachusetts General Hospital, Boston, MA, USA

SO J. Neurol. Sci. (1999), 171(1), 65-68

CODEN: JNSCAG; ISSN: 0022-510X

PB Elsevier Science Ireland Ltd.

DT Journal

LA English

RE.CNT 10

RE

(1) Boita, F; Sem Hosp Paris 1979, V55(9-10), P499 MEDLINE

(3) Daif, A; Stroke 1995, V26(7), P1193 MEDLINE

(4) Harper, P; Lancet 1988, V2(8617), P924 MEDLINE

(5) Hauser, D; Am J Ophthalmol 1996, V122(4), P592 MEDLINE

(6) Iranzo, A; J Neurol Neurosurg Psychiatry 1998, V64, P688 MEDLINE

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L32 ANSWER 3 OF 4 CA COPYRIGHT 2001 ACS

AN 132:263637 CA  
 TI Abnormalities in thrombin-antithrombin pathway in AL amyloidosis  
 AU Gamba, Gabriella; Montani, Nadia; Anesi, Ernesto; Palladini, Giovanni;  
 Lorenzutti, Federica; Perfetti, Vittorio; Merlini, Giampaolo  
 CS Department of Internal Medicine, University of Pavia, IRCCS Policlinico  
 San Matteo, Pavia, 27100, Italy  
 SO Amyloid (1999), 6(4), 273-277  
 CODEN: AIJIET; ISSN: 1350-6129  
 PB Parthenon Publishing Group  
 DT Journal  
 LA English  
 RE.CNT 25  
 RE  
 (10) Husby, G; Clin Immunol Immunopathol 1994, V70, P2 CA  
 (13) Khoory, M; J Clin Invest 1980, V65, P666 CA  
 (17) Marcatti, M; Thromb Res 1995, V80, P333 CA  
 (19) Merlini, G; Clin Chem 1981, V27, P1862 CA  
 (22) Sas, G; Thrombos Res 1975, V6, P87 CA  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L32 ANSWER 4 OF 4 CA COPYRIGHT 2001 ACS  
 AN 121:149109 CA  
 TI Treatment of neurodegenerative diseases with thrombin inhibitors  
 IN Friedrich, Thomas  
 PA BASF A.-G., Germany  
 SO Ger. Offen., 4 pp.  
 CODEN: GWXXBX  
 DT Patent  
 LA German  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 4303646	A1	19940811	DE 1993-4303646	19930209
	WO 9417821	A1	19940818	WO 1994-EP259	19940129
	W: CA, JP, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	CA 2153420	AA	19940818	CA 1994-2153420	19940129
	EP 683674	A1	19951129	EP 1994-906174	19940129
	R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, NL, PT, SE				
	JP 08507047	T2	19960730	JP 1994-517604	19940129
PRAI	DE 1993-4303646	19930209			
	WO 1994-EP259	19940129			

=> d 132 4 all

L32 ANSWER 4 OF 4 CA COPYRIGHT 2001 ACS  
 AN 121:149109 CA  
 TI Treatment of neurodegenerative diseases with thrombin inhibitors  
 IN Friedrich, Thomas  
 PA BASF A.-G., Germany  
 SO Ger. Offen., 4 pp.  
 CODEN: GWXXBX  
 DT Patent  
 LA German  
 IC ICM A61K037-02  
 CC 1-11 (Pharmacology)  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 4303646	A1	19940811	DE 1993-4303646	19930209
	WO 9417821	A1	19940818	WO 1994-EP259	19940129

W: CA, JP, US  
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE  
CA 2153420 AA 19940818 CA 1994-2153420 19940129  
EP 683674 A1 19951129 EP 1994-906174 19940129  
R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, NL, PT, SE  
JP 08507047 T2 19960730 JP 1994-517604 19940129  
PRAI DE 1993-4303646 19930209  
WO 1994-EP259 19940129  
AB Thrombin inhibitors, which may be combined with NGF, are useful in prepn. of medicaments for treatment of neurodegenerative diseases and disorders resulting e.g. from HIV-induced **neuropathy**, ischemia, subarachnoid hemorrhage, stroke, Alzheimer's disease, Huntington's disease, and parkinsonism. Thus, intracerebroventricular infusion of hirudin or protease nexin-1 (1-100 .mu.g/kg) into rats with unilateral brain lesions improved memory performance and the no. of choline acetyltransferase-pos. cortical neurons compared to those in operated but sham-treated rats.  
ST nerve degeneration treatment thrombin inhibitor; hirudin nerve degeneration treatment; proteinase nexin nerve degeneration treatment  
IT Memory, biological  
(after brain lesion, thrombin inhibitor effect on)  
IT Brain  
(regeneration of, after lesion, thrombin inhibitor effect on)  
IT Nerve, disease  
(degeneration, treatment of, with thrombin inhibitors)  
IT Brain, disease  
(lesion, nerve regeneration after, thrombin inhibitor effect on)  
IT 8001-27-2, Hirudin 9000-94-6, Antithrombin 148196-69-4  
RL: BIOL (Biological study)  
(nerve degeneration treatment with)  
IT 9061-61-4, Nerve growth factor  
RL: BIOL (Biological study)  
(nerve degeneration treatment with thrombin inhibitor and)

=> s 129 and 113

L33 0 L29 AND L13

=> s 129 and 112

L34 0 L29 AND L12

=> s 129 and 111

L35 30 L29 AND L11

=> d 135 15-30

L35 ANSWER 15 OF 30 CA COPYRIGHT 2001 ACS  
AN 119:220257 CA  
TI Properties of partly preinhibited hen brain **neuropathy** target esterase  
AU Vicedo, J. L.; Carrera, V.; Barril, J.; Vilanova, E.  
CS Dep. Neurochem., Alicante Univ., Alicante, Spain  
SO Chem.-Biol. Interact. (1993), 87(1-3), 417-23  
CODEN: CBINA8; ISSN: 0009-2797  
DT Journal  
LA English

L35 ANSWER 16 OF 30 CA COPYRIGHT 2001 ACS  
AN 119:153659 CA

TI Prophylaxis against and promotion of organophosphate-induced delayed neuropathy by phenyl di-n-pentylphosphinate  
AU Johnson, M. K.; Read, D. J.  
CS MRC Toxicol. Unit, Univ. Leicester, Leicester, LE1 9HN, UK  
SO Chem.-Biol. Interact. (1993), 87(1-3), 449-55  
CODEN: CBINA8; ISSN: 0009-2797  
DT Journal  
LA English

L35 ANSWER 17 OF 30 CA COPYRIGHT 2001 ACS  
AN 118:53794 CA  
TI Local application of neuropathic organophosphorus compounds to hen sciatic nerve: inhibition of neuropathy target esterase and peripheral neurological impairments  
AU Carrera, Victoria; Barril, Jose; Mauricio, Maricruz; Pellin, Maricruz; Vilanova, Eugenio  
CS Dep. Neurochem., Univ. Alicante, Alicante, 03002, Spain  
SO Toxicol. Appl. Pharmacol. (1992), 117(2), 218-25  
CODEN: TXAPA9; ISSN: 0041-008X  
DT Journal  
LA English

L35 ANSWER 18 OF 30 CA COPYRIGHT 2001 ACS  
AN 118:34045 CA  
TI Phenylmethanesulfonyl fluoride elicits and intensifies the clinical expression of neuropathic insults  
AU Moretto, A.; Bertolazzi, M.; Capodicasa, E.; Peraica, M.; Richardson, R. J.; Scapellato, M. L.; Lotti, M.  
CS Ist. Med. Lavoro, Univ. Stud. Padova, Padua, I-35127, Italy  
SO Arch. Toxicol. (1992), 66(1), 67-72  
CODEN: ARTODN; ISSN: 0340-5761  
DT Journal  
LA English

L35 ANSWER 19 OF 30 CA COPYRIGHT 2001 ACS  
AN 118:2118 CA  
TI Clinical expression of organophosphate-induced delayed polyneuropathy in rats  
AU Moretto, Angelo; Capodicasa, Eugenio; Lotti, Marcello  
CS Ist. Med. Lavoro, Univ. Padova, Padua, 35127, Italy  
SO Toxicol. Lett. (1992), 63(1), 97-102  
CODEN: TOLED5; ISSN: 0378-4274  
DT Journal  
LA English

L35 ANSWER 20 OF 30 CA COPYRIGHT 2001 ACS  
AN 117:165625 CA  
TI The inhibitory effect of neuropathic organophosphate esters on neurite outgrowth in cell cultures: a basis for screening for delayed neurotoxicity  
AU Henschler, D.; Schmuck, G.; Van Aerssen, M.; Schiffmann, D.  
CS Inst. Toxicol., Univ. Wuerzburg, Wuerzburg, D-8700, Germany  
SO Toxicol. in Vitro (1992), 6(4), 327-35  
CODEN: TIVIEQ; ISSN: 0887-2333  
DT Journal  
LA English

L35 ANSWER 21 OF 30 CA COPYRIGHT 2001 ACS  
AN 115:176927 CA  
TI Promotion of organophosphates induced delayed polyneuropathy by phenylmethanesulfonyl fluoride. Comments  
AU Pope, Carey N.; Padilla, Stephanie

CS Sch. Pharm., Northeast Louisiana Univ., Monroe, LA, 71209, USA  
SO Toxicol. Appl. Pharmacol. (1991), 110(1), 179-80  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English

L35 ANSWER 22 OF 30 CA COPYRIGHT 2001 ACS  
AN 115:43729 CA  
TI Promotion of organophosphate-induced delayed polyneuropathy by phenylmethanesulfonyl fluoride  
AU Lotti, Marcello; Caroldi, Stefano; Capodicasa, Eugenio; Moretto, Angelo  
CS Ist. Med. Lav., Univ. Padova, Padua, I-35127, Italy  
SO Toxicol. Appl. Pharmacol. (1991), 108(2), 234-41  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English

L35 ANSWER 23 OF 30 CA COPYRIGHT 2001 ACS  
AN 110:149296 CA  
TI Triphenyl phosphite neurotoxicity in the hen: inhibition of neurotoxic esterase and of prophylaxis by phenylmethanesulfonyl fluoride  
AU Carrington, Clark D.; Abou-Donia, Mohamed B.  
CS Med. Cent., Duke Univ., Durham, NC, 27710, USA  
SO Arch. Toxicol. (1988), 62(5), 375-80  
CODEN: ARTODN; ISSN: 0340-5761  
DT Journal  
LA English

L35 ANSWER 24 OF 30 CA COPYRIGHT 2001 ACS  
AN 106:190736 CA  
TI Central-peripheral delayed **neuropathy** caused by diisopropyl phosphorofluoridate (DFP): segregation of peripheral nerve and spinal cord effects using biochemical, clinical, and morphological criteria  
AU Lotti, M.; Caroldi, S.; Moretto, A.; Johnson, M. K.; Fish, C. J.; Gopinath, C.; Roberts, N. L.  
CS Ist. Med. Lavoro, Univ. Padova, Padua, 35127, Italy  
SO Toxicol. Appl. Pharmacol. (1987), 88(1), 87-96  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English

L35 ANSWER 25 OF 30 CA COPYRIGHT 2001 ACS  
AN 104:16305 CA  
TI Phenylmethanesulfonyl fluoride protects rats from mipafox-induced delayed **neuropathy**  
AU Veronesi, Bellina; Padilla, Stephanie  
CS Health Effects Res. Lab., U. S. Environ. Prot. Agency, Research Triangle Park, NC, 27711, USA  
SO Toxicol. Appl. Pharmacol. (1985), 81(2), 258-64  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English

L35 ANSWER 26 OF 30 CA COPYRIGHT 2001 ACS  
AN 102:107664 CA  
TI Neurotoxic esterase in fooster testis  
AU Lotti, Marcello; Wei, Eddie T.; Spear, Robert C.; Becker, Charles E.  
CS North. California Occup. Health Cent., Univ. California, San Francisco, CA, USA  
SO Toxicol. Appl. Pharmacol. (1985), 77(1), 175-80  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal

LA English

L35 ANSWER 27 OF 30 CA COPYRIGHT 2001 ACS  
AN 102:1635 CA  
TI Intraarterial injection of diisopropylfluorophosphate or phenylmethanesulfonyl fluoride produces unilateral **neuropathy** or protection, respectively, in hens  
AU Caroldi, Stefano; Lotti, Marcello; Masutti, Alberto  
CS Ist. Med. Lavoro, Univ. Padova, Padua, 35127, Italy  
SO Biochem. Pharmacol. (1984), 33(20), 3213-17  
CODEN: BCPCA6; ISSN: 0006-2952  
DT Journal  
LA English

L35 ANSWER 28 OF 30 CA COPYRIGHT 2001 ACS  
AN 99:207626 CA  
TI An electrophysiologic and ultrastructural study of the phenylmethanesulfonyl fluoride protection against a delayed organophosphorus **neuropathy**  
AU Drakontides, Anna B.; Baker, Thomas  
CS Dep. Anat., New York Med. Coll., Valhalla, NY, 10595, USA  
SO Toxicol. Appl. Pharmacol. (1983), 70(3), 411-22  
CODEN: TXAPA9; ISSN: 0041-008X  
DT Journal  
LA English

L35 ANSWER 29 OF 30 CA COPYRIGHT 2001 ACS  
AN 94:97371 CA  
TI The effects of phenylmethanesulfonyl fluoride on delayed organophosphorus **neuropathy**  
AU Baker, Thomas; Lowndes, Herbert E.; Johnson, Martin K.; Sandborg, Irene C.  
CS Med. Coll., Cornell Univ., New York, NY, 10021, USA  
SO Arch. Toxicol. (1980), 46(3-4), 305-11  
CODEN: ARTODN; ISSN: 0340-5761  
DT Journal  
LA English

L35 ANSWER 30 OF 30 CA COPYRIGHT 2001 ACS  
AN 91:69597 CA  
TI Neurotoxicity of organophosphorus pesticides: predictions can be based on in vitro studies with hen and human enzymes  
AU Lotti, Marcello; Johnson, Martin Keith  
CS Mol. Toxicol. Sect., MRC, Carshalton/Surrey, SM5 4EF, Engl.  
SO Arch. Toxicol. (1978), 41(3), 215-21  
CODEN: ARTODN; ISSN: 0340-5761  
DT Journal  
LA English

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1 11 S NEUROSERPIN  
L2 1 S PMSF  
L3 1 S APMSF  
L4 1 S L3  
L5 5 S ANTIPAIN  
L6 170 S ANTITHROMBIN

L7 11 S LEUPEPTIN  
L8 9 S DICHLOROCOUMARIN  
L9 2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

L10 17 S L1  
E NEUROSERPIN  
L11 728 S L2 → PMSF  
L12 24 S L3  
L13 343 S L5  
L14 4326 S L6  
L15 171 S L7  
L16 16 S L8  
L17 372 S L9  
E NEUROPATHY  
L18 5490 S E3  
E EPILEPSY  
L19 10805 S E3-E12  
E SEIZURE  
L20 14562 S E3-E6  
E HYPOXIA  
L21 24801 S E3  
E STROKE  
L22 11880 S E3  
L23 0 S L17 AND L18  
L24 0 S L17 AND L19  
L25 0 S L17 AND L20  
L26 0 S L17 AND L21  
L27 0 S L17 AND L22  
L28 0 S L16 AND L18  
L29 25960 S L18 OR L19 OR L20  
L30 0 S L29 AND L16  
L31 0 S L29 AND L15  
L32 4 S L29 AND L14  
L33 0 S L29 AND L13  
L34 0 S L29 AND L12  
L35 30 S L29 AND L11

=> d 135 7-14

L35 ANSWER 7 OF 30 CA COPYRIGHT 2001 ACS

AN 127:216150 CA

TI Phenyl valerate esterases other than **neuropathy** target esterase  
and the promotion of organophosphate polyneuropathy

AU Milatovic, Dejan; Moretto, Angelo; Osman, Khaled A.; Lotti, Marcello

CS Istituto di Medicina del Lavoro, Universita degli Studi di Padova, Padua,  
I-35127, Italy

SO Chem. Res. Toxicol. (1997), 10(9), 1045-1048

CODEN: CRTOEC; ISSN: 0893-228X

PB American Chemical Society

DT Journal

LA English

L35 ANSWER 8 OF 30 CA COPYRIGHT 2001 ACS

AN 127:215998 CA

TI Improved in vitro method for screening organophosphate-induced delayed  
polyneuropathy

AU Schmuck, G.; Ahr, H. J.

CS Bayer AG, Pharma Research Centre, Wuppertal, D-42096, Germany

SO Toxicol. in Vitro (1997), 11(3), 263-270

CODEN: TIVIEQ; ISSN: 0887-2333

PB Elsevier

DT Journal

LA English

L35 ANSWER 9 OF 30 CA COPYRIGHT 2001 ACS

AN 125:294975 CA

TI Sulfonyl fluorides and the promotion of diisopropyl fluorophosphate **neuropathy**

AU Osman, Khaled A.; Moretto, Angelo; Lotti, Marcello

CS Instituto di Medicina del Lavoro, Universita degli Studi di Padova, Padua,

35127, Italy

SO Fundam. Appl. Toxicol. (1996), 33(2), 294-297

CODEN: FAATDF; ISSN: 0272-0590

DT Journal

LA English

L35 ANSWER 10 OF 30 CA COPYRIGHT 2001 ACS

AN 125:134966 CA

TI Subacute neurotoxicity induced in mice by potent organophosphorus **neuropathy** target esterase inhibitors

AU Wu, Shao-Yong; Casida, John E.

CS Environmental Chemistry and Toxicology Lab., Univ. of California, Berkeley, CA, 94720-3112, USA

SO Toxicol. Appl. Pharmacol. (1996), 139(1), 195-202

CODEN: TXAPAA9; ISSN: 0041-008X

DT Journal

LA English

L35 ANSWER 11 OF 30 CA COPYRIGHT 2001 ACS

AN 124:223242 CA

TI Effects of various post-treatment by phenylmethylsulfonyl fluoride on delayed neurotoxicity induced by leptophos

AU Piao, Feng Yuan; Kitabatake, Masayoshi; Xie, Xiu Kui; Yamauchi, Toru

CS School Medicine, Mie University, Edobashi, 514, Japan

SO J. Toxicol. Sci. (1995), 20(5), 609-17

CODEN: JTSCDR; ISSN: 0388-1350

DT Journal

LA English

L35 ANSWER 12 OF 30 CA COPYRIGHT 2001 ACS

AN 124:2746 CA

TI Triphenylphosphite **neuropathy** in hens

AU Fioroni, F.; Moretto, A.; Lotti, M.

CS Ist. Med. Lavoro, Univ. Studi Padova, Padua, I-35127, Italy

SO Arch. Toxicol. (1995), 69(10), 705-11

CODEN: ARTODN; ISSN: 0340-5761

DT Journal

LA English

L35 ANSWER 13 OF 30 CA COPYRIGHT 2001 ACS

AN 123:332411 CA

TI Selective promotion by phenylmethanesulfonyl fluoride of peripheral and spinal cord neuropathies initiated by diisopropyl phosphorofluoridate in the hen

AU Peraica, Maja; Moretto, Angelo; Lotti, Marcello

CS Universita degli Studi di Padova, Istituto di Medicina del Lavoro, Via Faccioliati 71, Padua, 35127, Italy

SO Toxicol. Lett. (1995), 80(1-3), 115-21

CODEN: TOLED5; ISSN: 0378-4274

DT Journal

LA English

L35 ANSWER 14 OF 30 CA COPYRIGHT 2001 ACS  
AN 123:104746 CA  
TI Effect of organophosphorus compounds on **neuropathy** target esterase in hens  
AU Sadek, Omayma M.; Abdelhamid, Essam E.; El-Sayed, Mohamed M.;  
Abdel-Moneam, Nehad M.; Mansour, Nabil A.  
CS Faculty of Science, Alexandria University, Egypt  
SO Egypt. J. Biochem. (1995), 13(1), 143-52  
CODEN: EGJBE4; ISSN: 1012-554X  
DT Journal  
LA English

=> d 135 25 23 16 all

L35 ANSWER 25 OF 30 CA COPYRIGHT 2001 ACS  
AN 104:16305 CA  
TI Phenylmethylsulfonyl fluoride protects rats from mipafox-induced delayed **neuropathy**  
AU Veronesi, Bellina; Padilla, Stephanie  
CS Health Effects Res. Lab., U. S. Environ. Prot. Agency, Research Triangle Park, NC, 27711, USA  
SO Toxicol. Appl. Pharmacol. (1985), 81(2), 258-64  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English  
CC 4-4 (Toxicology)  
AB Prior exposure to a nonaging **neuropathy** target enzyme (NTE) inhibitor, phenylmethylsulfonyl fluoride (PMSF) [329-98-6], protects rats from neurol. damage after subsequent exposure to mipafox [371-86-8]. Adult, male rats were exposed to either PMSF (250 mg/kg, s.c.) or to mipafox (15 mg/kg, i.p.) and a time course of brain NTE inhibition and recovery was defined. A sep. group of PMSF-treated rats was exposed to mipafox when brain NTE inhibition was 87.7%. Conversely, another group of rats, pretreated with mipafox, was dosed with PMSF when NTE inhibition was 90.2%. A 3rd group of animals, treated with PMSF, was exposed to mipafox 14 days later, when NTE activity had recovered to within 10% of control amts. Histopathol. survey (14-21 days postexposure) indicated severed cervical cord damage (damage score .gtoreq.3) in the following frequencies: PMSF, 0%; mipafox, 85%; PMSF-4 h-mipafox, 0%; mipafox-4 h-PMSF, 100%; PMSF-14 days-mipafox, 75%; controls, 0%. These data indicate that PMSF pretreatment protects rats against mipafox-induced neurol. damage and that the timing of administration and order of presentation are crit. to this protection. Apparently, the initiation of organophosphorus-induced delayed **neuropathy** is a multistage event involving inhibition and aging, and these stages are exptl. separable.  
ST mipafox neurotoxicity phenylmethylsulfonyl fluoride  
IT Brain, composition  
(**neuropathy** target enzyme of, mipafox effect on,  
phenylmethylsulfonyl fluoride in relation to)  
IT Spinal cord  
(cervical, mipafox toxicity to, phenylmethylsulfonyl fluoride  
protection against)  
IT 329-98-6  
RL: BIOL (Biological study)  
(mipafox neurotoxicity protection by)  
IT 9013-79-0  
RL: BIOL (Biological study)  
(neurotoxic, of brain, mipafox effect on, phenylmethylsulfonyl fluoride)

in relation to)

IT 371-86-8

RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)  
(neurotoxicity of, phenylmethylsulfonyl fluoride protection against)

L35 ANSWER 23 OF 30 CA COPYRIGHT 2001 ACS

AN 110:149296 CA

TI Triphenyl phosphite neurotoxicity in the hen: inhibition of neurotoxic esterase and of prophylaxis by phenylmethylsulfonyl fluoride

AU Carrington, Clark D.; Abou-Donia, Mohamed B.

CS Med. Cent., Duke Univ., Durham, NC, 27710, USA

SO Arch. Toxicol. (1988), 62(5), 375-80

CODEN: ARTODN; ISSN: 0340-5761

DT Journal

LA English

CC 4-3 (Toxicology)

AB The neuropathic syndrome resulting in the cat and the rat from single or multiple doses of the phosphorous acid ester tri-Ph phosphite (TPP) has been reported to differ from the syndrome caused by numerous phosphoric acid esters, which is known as organophosphorous compd.-induced delayed neurotoxicity (OPIDN). Since the hen is used to test compds. for OPIDN, the neurotoxicity of single s.c. doses of TPP was studied using this animal model. TPP (1000 mg/kg) produced progressive ataxia and paralysis which began to develop 5-10 days after dosing. Similar signs were obsd. when s.c. doses of the OPIDN-causing agents tri-o-cresyl phosphate (TOCP) or diisopropyl phosphorofluoridate (DFP) were administered. The min. neurotoxic dose of TPP was 500 mg/kg. Prior administration of phenylmethylsulfonyl fluoride (PMSF) prevented the development of a neuropathy induced by DFP, but did not fully protect the hens from TPP or TOCP. PMSF slowed, but did not prevent, the neuropathy caused by TOCP. PMSF reduced the neurotoxicity of 500 mg/kg TPP, but increased the neurotoxicity of 1000 mg/kg TPP. TPP was a very potent inhibitor of neurotoxic esterase (NTE), the putative target site for OPIDN, in vitro, with a ki of .apprx.2.1 .times. 105 M-1 min-1.

Equimolar

doses of either TPP (1000 mg/kg) and TOCP (1187 mg/kg) caused over 80% inhibition of neurotoxic esterase (NTE) in brain and sciatic nerve. This high level of NTE inhibition persisted for several weeks. This prolonged inhibition probably accounts for the inability of PMSF to block the neurotoxicity of TOCP. The dose-response curve for NTE inhibition 48 h after dosing indicated that a level of 70% inhibition correlated with the neurotoxicity of TPP. Subneurotoxic doses of TPP and DFP had an additive effect which could be blocked by PMSF. These results indicate that TPP can cause OPIDN in the hen. The synergism between PMSF and the higher dose of TPP suggests the presence of a 2nd neurotoxic effect as well.

ST triphenyl phosphite neurotoxicity chicken phenylmethylsulfonyl fluoride; neurotoxic esterase triphenyl phosphite chicken

IT Paralysis

(from tri-Ph phosphite, in hen, phenylmethylsulfonyl fluoride effect on)

IT Brain, composition

(neurotoxic esterase of, of hen, tri-Ph phosphite effect on)

IT Chicken

(tri-Ph phosphite neurotoxicity in, phenylmethylsulfonyl fluoride effect on)

IT Nervous system

(disease, ataxia, from tri-Ph phosphite, in hen, phenylmethylsulfonyl fluoride effect on)

IT Nerve, toxic chemical and physical damage

(neuropathy, from tri-Ph phosphite, in hen, phenylmethylsulfonyl fluoride effect on)

IT Organic compounds, biological studies

RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)

(phosphorus-contg., neurotoxicity of, in hen, phenylmethylsulfonyl fluoride effect on)  
IT Nerve, composition  
(sciatic, neurotoxic esterase of, of hen, tri-Ph phosphite effect on)  
IT 9013-79-0, Esterase  
RL: BIOL (Biological study)  
(neurotoxic, of brain and sciatic nerve of hen, tri-Ph phosphite effect on)  
on)  
IT 101-02-0, Triphenyl phosphite  
RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)  
(neurotoxicity of, in hen, phenylmethylsulfonyl fluoride effect on)  
IT 55-91-4, DFP 78-30-8, TOCP  
RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)  
(neurotoxicity of, in hen, phenylmethylsulfonyl fluoride effect on,  
tri-Ph phosphite in relation to)  
IT 329-98-6, Phenylmethylsulfonyl fluoride  
RL: BIOL (Biological study)  
(triphenylphosphite neurotoxicity response to, in hen)

L35 ANSWER 16 OF 30 CA COPYRIGHT 2001 ACS

AN 119:153659 CA

TI Prophylaxis against and promotion of organophosphate-induced delayed neuropathy by phenyl di-n-pentylphosphinate

AU Johnson, M. K.; Read, D. J.

CS MRC Toxicol. Unit, Univ. Leicester, Leicester, LE1 9HN, UK

SO Chem.-Biol. Interact. (1993), 87(1-3), 449-55

CODEN: CBINA8; ISSN: 0009-2797

DT Journal

LA English

CC 4-3 (Toxicology)

AB Ph di-n-pentylphosphinate (PPP) is a potent inhibitor of neuropathy target esterase (NTE) with a negligible effect on acetylcholinesterase; I50s at 37.degree.C for 20 min and pH 8, resp. are 0.2 .mu.M and >2mM. PPP is not neuropathic. This is compatible with the fact that inhibited NTE in autopsy material from hens dosed with PPP can always be reactivated in vitro, presumably because no 'aging' reaction has

occurred. PPP (10 mg/kg s.c.) given to hens up to 4 days before severely neuropathic doses (1.7 mg/kg) of diisopropylphosphorofluoridate (DFP) prevented neuropathic but not cholinergic effects of DFP. Hens given PPP 3 days after a sub-neuropathic dose of DFP (0.4 mg/kg) developed severe clin. neuropathy (clin. scores of 7 and 5 compared with DFP-plus-solvent scores 0, 1, 3). These prophylactic and promoting effects are similar to those exerted by phenylmethanesulfonyl fluoride (PMSF) at doses which inhibit NTE. In 3 out of 4 birds, a pre-dose with PMSF (15 mg/kg) prevented the promoting effect of 120 mg/kg PMSF given after DFP.

ST phenyl dipentylphosphinate neuropathy

IT Nerve, disease

(neuropathy, from Ph dipentylphosphinate, prophylaxis against)

IT 55-91-4, Diisopropylphosphorofluoridate 329-98-6,

Phenylmethanesulfonyl fluoride

RL: BIOL (Biological study)

(neuropathy by Ph dipentylphosphonate prevention by)

IT 14656-17-8

RL: BIOL (Biological study)

(neuropathy from, prophylaxis against)

=> s 129 and 110

L36 O L29 AND L10

=> s l21 and l22

L37 350 L21 AND L22

=> s l21 or l22

L38 36331 L21 OR L22

=> s l38 and l10

L39 4 L38 AND L10

=> d l39 1-4

L39 ANSWER 1 OF 4 CA COPYRIGHT 2001 ACS

AN 133:248681 CA

TI Human brain-associated inhibitor of tissue-type plasminogen activator (BAIT) and cDNA and their use for treatment of **stroke**

IN Lawrence, Daniel A.; Yepes, Manuel; Sandkvist, Maria; Coleman, Timothy A.;

Wong, Michael K. K.

PA Human Genome Sciences, Inc., USA; American Red Cross

SO PCT Int. Appl., 302 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000053793	A1	20000914	WO 2000-US5956	20000308
	W:	AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,			
TM	RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			

PRAI US 1999-123704 19990310

RE.CNT 1

RE

(1) Hastings; US 6008020 1999 CA

L39 ANSWER 2 OF 4 CA COPYRIGHT 2001 ACS

AN 133:232282 CA

TI Serine protease inhibitors: novel therapeutic targets for **stroke** ?

AU Vivien, Denis; Buisson, Alain

CS Universite de Caen, Caen, 14074, Fr.

SO J. Cereb. Blood Flow Metab. (2000), 20(5), 755-764

CODEN: JCBMDN; ISSN: 0271-678X

PB Lippincott Williams & Wilkins

DT Journal; General Review

LA English

RE.CNT 69

RE

(2) Baranes, D; Neuron 1998, V21, P813 CA

(3) Berger, P; Gene 1998, V214, P25 CA

(4) Buisson, A; FASEB J 1998, V12, P1683 CA

(6) Carmeliet, P; Nature 1994, V368, P419 CA

(7) Chen, Z; Cell 1997, V91, P917 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L39 ANSWER 3 OF 4 CA COPYRIGHT 2001 ACS

AN 133:187780 CA

TI Neuroserpin reduces cerebral infarct volume and protects neurons from ischemia-induced apoptosis

AU Yepes, Manuel; Sandkvist, Maria; Wong, Mike K. K.; Coleman, Timothy A.; Smith, Elizabeth; Cohan, Stanley L.; Lawrence, Daniel A.

CS Department of Biochemistry, American Red Cross Holland Laboratory, Rockville, MD, 20855, USA

SO Blood (2000), 96(2), 569-576

CODEN: BLOOAW; ISSN: 0006-4971

PB American Society of Hematology

DT Journal

LA English

RE.CNT 70

RE

(1) Ahn, M; Brain Res 1999, V837, P169 CA

(3) Benveniste, H; J Neurochem 1984, V43, P1369 CA

(5) Calof, A; Neuron 1994, V13, P117 CA

(7) Carroll, P; Development 1994, V120, P3173 CA

(8) Chen, Z; Cell 1997, V91, P917 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L39 ANSWER 4 OF 4 CA COPYRIGHT 2001 ACS

AN 131:139510 CA

TI Neuroserpin applications as a pharmaceutical or diagnostic agent

IN Sonderegger, Peter; Schrimpf, Sabine Petra; Kruger, Stefan Robert; Osterwalder, Thomas; Stockli, Esther Trudi

PA Switz.

SO PCT Int. Appl., 55 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9941381	A1	19990819	WO 1999-IB248	19990212
	W:	AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,			
TM	RW:	GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			

AU 9921807 A1 19990830 AU 1999-21807 19990212

PRAI US 1998-23129 19980213

WO 1999-IB248 19990212

RE.CNT 6

RE

(1) Coleman, T; WO 9816643 A 1998 CA

(2) Hastings, G; THE JOURNAL OF BIOLOGICAL CHEMISTRY 1997, V272(52), P33062 CA

(3) Incyte Pharma Inc; WO 9640922 A 1996 CA

(4) Krueger, S; THE JOURNAL OF NEUROSCIENCE 1997, V17(23), P8984 CA

(6) Schrimpf; GENOMICS 1997, V40(1), P55 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d 139 4 all

L39 ANSWER 4 OF 4 CA COPYRIGHT 2001 ACS  
 AN 131:139510 CA  
 TI Neuroserpin applications as a pharmaceutical or diagnostic agent  
 IN Sonderegger, Peter; Schrimpf, Sabine Petra; Kruger, Stefan Robert;  
 Osterwalder, Thomas; Stockli, Esther Trudi  
 PA Switz.  
 SO PCT Int. Appl., 55 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 IC ICM C12N015-15  
 ICS C07K014-81; A61K038-17; A61K048-00; A01K067-027  
 CC 1-11 (Pharmacology)  
 Section cross-reference(s): 3, 7

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9941381	A1	19990819	WO 1999-IB248	19990212
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,				

TM	RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	AU 9921807	A1	19990830	AU 1999-21807	19990212

PRAI US 1998-23129 19980213  
 WO 1999-IB248 19990212

AB Pharmaceutical and diagnostic applications of neuroserpins, in particular human neuroserpin, are provided. Neuroserpin expression is enhanced in neurons of the ipsilateral hemisphere after focal ischemia **stroke**. In the adult brain, neuroserpin and tissue-type plasminogen activator (tPA) for complexes. Overexpression of neuroserpin in central nervous system neurons using transgenic mice technol. results in reduced tPA activity in the brain and an attenuated microglial activation in the reactive zone of a focal ischemic **stroke**. Thus, neuroserpins are valuable agents in the treatment of disorders of the nervous system, in particular the central nervous system. They are very useful in the treatment of **stroke** and for the development of drugs.

ST neuroserpin pharmacol diagnosis nervous system; sequence neuroserpin cDNA human mouse; drug screening neuroserpin nervous system; **stroke** treatment neuroserpin

IT Diagnosis  
 (agents; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Antitumor agents  
 (brain; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Nervous system  
 (central, disease, treatment of; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Neuron  
 (death, prevention of; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Nervous system  
 (disease, treatment of; neuroserpin applications as a pharmaceutical or  
 or  
 diagnostic agent)

IT cDNA sequences

(for neuroserpin from human and mouse)

IT Brain, neoplasm  
(inhibitors; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Blood vessel, neoplasm  
(metastasis inhibitors; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Antitumor agents

Drug screening

Drugs

Molecular cloning

Mouse  
(neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Protein sequences  
(of neuroserpin from human and mouse)

IT Antibodies

Antigens

RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP (Preparation)  
(prodn. of; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Escherichia coli

Eukaryote (Eukaryotae)  
(recombinant expression host; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Brain, disease  
(stroke, treatment or prevention of tissue destruction in; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Animal  
(transgenic; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT Angiogenesis

Brain, disease  
(treatment of; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT 188310-87-4, Neuroserpin (human gene PI12 precursor)  
188364-82-1, Neuroserpin 200890-63-7, Neuroserpin (mouse strain BALB/c brain)  
RL: ARU (Analytical role, unclassified); BAC (Biological activity or effector, except adverse); PRP (Properties); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)  
(amino acid sequence; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT 9001-12-1, Collagenase 9001-90-5, Plasmin 9004-06-2, Elastase 9025-26-7, Cathepsin D 9032-92-2, Glycosidase 9039-53-6, Urokinase 9040-48-6, Gelatinase 9047-22-7, Cathepsin B 37353-41-6, Cysteine proteinase 56645-49-9, Cathepsin G 79955-99-0, Stromelysin 139639-23-9, Tissue-type plasminogen activator 139639-24-0, Urokinase-type plasminogen activator 141256-52-2, Matrilysin 141907-41-7, Matrix metalloproteinase  
RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
(nervous system disorders requiring inhibition of; neuroserpin applications as a pharmaceutical or diagnostic agent)

IT 185376-12-9, GenBank Z81326 197679-81-5, GenBank AJ001700  
RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
(nucleotide sequence; neuroserpin applications as a pharmaceutical or diagnostic agent)

RE.CNT 6

RE

- (1) Coleman, T; WO 9816643 A 1998 CA
- (2) Hastings, G; THE JOURNAL OF BIOLOGICAL CHEMISTRY 1997, V272(52), P33062 CA
- (3) Incyte Pharma Inc; WO 9640922 A 1996 CA
- (4) Krueger, S; THE JOURNAL OF NEUROSCIENCE 1997, V17(23), P8984 CA
- (5) Marra, M; EMBL DATABASE ENTRY
- (6) Schrimpf; GENOMICS 1997, V40(1), P55 CA

=> s 138 and l11

L40 2 L38 AND L11

=> d 140 1-2

L40 ANSWER 1 OF 2 CA COPYRIGHT 2001 ACS

AN 120:49588 CA

TI Method for processing and preserving collagen-based tissues for transplantation

IN Livesey, Stephen A.; Del Campo, Anthony A.; Nag, Abhijit; Nichols, Ken B.;

Griffey, Edward S.; Coleman, Christopher

PA Lifecell Corp., USA

SO Can. Pat. Appl., 63 pp.

CODEN: CPXXEB

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	CA 2089336	AA	19930813	CA 1993-2089336	19930211
	CA 2051092	AA	19920313	CA 1991-2051092	19910910
	AU 9183797	A1	19920319	AU 1991-83797	19910910
	AU 650045	B2	19940609		
	EP 475409	A2	19920318	EP 1991-115480	19910912
	EP 475409	A3	19930901		
	EP 475409	B1	19980415		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE				
	AT 164981	E	19980515	AT 1991-115480	19910912
	ES 2114868	T3	19980616	ES 1991-115480	19910912
	US 5336616	A	19940809	US 1993-4752	19930202
	AU 9332934	A1	19930819	AU 1993-32934	19930210
	AU 668703	B2	19960516		
	EP 564786	A2	19931013	EP 1993-102264	19930212
	EP 564786	A3	19940706		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT,				
SE	JP 06261933	A2	19940920	JP 1993-47373	19930212
	US 5364756	A	19941115	US 1993-18357	19930216
	AU 9467405	A1	19940922	AU 1994-67405	19940713
	AU 677845	B2	19970508		
	US 5780295	A	19980714	US 1996-752740	19961114
	US 6194136	B1	20010227	US 1998-114433	19980713
PRAI	US 1992-835138	19920212			
	US 1993-4752	19930202			
	US 1990-581584	19900912			
	US 1991-709504	19910603			
	US 1993-18357	19930216			
	US 1994-291340	19940817			
	US 1996-18357	19960216			
	US 1996-752740	19961114			

L40 ANSWER 2 OF 2 CA COPYRIGHT 2001 ACS  
AN 117:103472 CA  
TI A novel cytotoxicity screening assay using a multiwell fluorescence scanner  
AU Nieminen, Anna Liisa; Gores, Gregory J.; Bond, John M.; Imberti, Roberto; Herman, Brian; Lemasters, John J.  
CS Sch. Med., Univ. North Carolina, Chapel Hill, NC, 27599-7090, USA  
SO Toxicol. Appl. Pharmacol. (1992), 115(2), 147-55  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English

=> s 138 and 112

L41 0 L38 AND L12

=> s 138 and 113

L42 1 L38 AND L13

=> d 142

L42 ANSWER 1 OF 1 CA COPYRIGHT 2001 ACS  
AN 117:103472 CA  
TI A novel cytotoxicity screening assay using a multiwell fluorescence scanner  
AU Nieminen, Anna Liisa; Gores, Gregory J.; Bond, John M.; Imberti, Roberto; Herman, Brian; Lemasters, John J.  
CS Sch. Med., Univ. North Carolina, Chapel Hill, NC, 27599-7090, USA  
SO Toxicol. Appl. Pharmacol. (1992), 115(2), 147-55  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English

=> d 142 1 all

L42 ANSWER 1 OF 1 CA COPYRIGHT 2001 ACS  
AN 117:103472 CA  
TI A novel cytotoxicity screening assay using a multiwell fluorescence scanner  
AU Nieminen, Anna Liisa; Gores, Gregory J.; Bond, John M.; Imberti, Roberto; Herman, Brian; Lemasters, John J.  
CS Sch. Med., Univ. North Carolina, Chapel Hill, NC, 27599-7090, USA  
SO Toxicol. Appl. Pharmacol. (1992), 115(2), 147-55  
CODEN: TXAPAA9; ISSN: 0041-008X  
DT Journal  
LA English  
CC 1-1 (Pharmacology)  
AB A new assay using a multiwell fluorescence scanner was developed for screening cytotoxicity to cells cultured in 96-well microtiter plates. The assay is based on binding of propidium iodide to nuclei of cells whose plasma membranes have become permeable due to cell death. Fluorescence of propidium iodide measured with a multiwell fluorescence scanner increased in proportion to the no. of permeabilized cells. After ATP depletion of hepatocytes and neonatal cardiac myocytes with metabolic inhibitors ("chem. hypoxia"), and exposure of Madine Darby canine kidney cells to the toxic chem., HgCl<sub>2</sub>, propidium iodide fluorescence

progressively increased. Increases of fluorescence were linearly proportional with release of lactate dehydrogenase into the culture medium. Employing this cytotoxicity screening assay, protection by various agents against lethal injury was evaluated in cultured hepatocytes

during chem. hypoxia. Inhibitors of cysteine proteases (i.e., antipain, leupeptin, E-64), serine proteases (i.e., PMSF), and aspartic acid proteases (i.e., pepstatin A) did not protect against chem.

hypoxia. In contrast, 1,10-phenanthroline, an inhibitor of metalloprotease, markedly protected against the onset of cell death

during

chem. hypoxia. Half-maximal protection after 60 min occurred at 0.5 .mu.M. Phospholipase inhibitors, chlorpromazine (50 .mu.M) and mepacrine (50 .mu.M), also substantially retarded cell killing. U74006F, an inhibitor of lipid peroxidn., slowed cell killing to a lesser extent during chem. hypoxia and after oxidative stress with tert-Bu hydroperoxide. Calciphor, a dimer of prostaglandin B1, did not protect against cell killing during chem. hypoxia or tert-Bu hydroperoxide toxicity. In conclusion, this high capacity cytotoxicity assay for cells cultured in 96-well microtiter plates is suitable for rapid screening of potential cytoprotective agents in a variety of cell types.

ST cytotoxicity screening assay cytoprotectant hepatocyte cardiomyocyte

IT Bioassay

(cytoprotectants screening by fluorescence, in hepatocyte and cardiomyocyte monolayers)

IT Heart

(cytoprotectants screening in monolayer culture of, using multiwell fluorescence scanner)

IT Leupeptins

RL: PRP (Properties)

(cytoprotective effect of, in cytotoxicity screening assay using multiwell fluorescence scanner)

IT Fluorescence

(cytotoxicity screening assay based on, hepatocyte and cardiomyocyte monolayers in)

IT Cytoprotective agents

(screening of, in cytotoxicity assay using multiwell fluorescence scanner)

IT Toxicity

(cytotoxicity, screening assay using multiwell fluorescence scanner, in hepatocyte and cardiomyocyte monolayers)

IT Liver

(hepatocyte, cytoprotectants screening in monolayer culture of, using multiwell fluorescence scanner)

IT 50-53-3, Chlorpromazine, biological studies 66-71-7,

1,10-Phenanthroline

83-89-6, Mepacrine 329-98-6, PMSF 26305-03-3, Pepstatin A

37691-11-5, Antipain 66701-25-5, E-64 110101-67-2

135608-06-9

RL: PRP (Properties)

(cytoprotective effect of, in cytotoxicity screening assay using multiwell fluorescence scanner)

=> d 138 and 13

'AND' IS NOT A VALID FORMAT FOR FILE 'CA'

The following are valid formats:

ABS ----- GI and AB

ALL ----- BIB, AB, IND, RE  
 APPS ----- AI, PRAI  
 BIB ----- AN, plus Bibliographic Data and PI table (default)  
 CAN ----- List of CA abstract numbers without answer numbers  
 CBIB ----- AN, plus Compressed Bibliographic Data  
 DALL ----- ALL, delimited (end of each field identified)  
 DMAX ----- MAX, delimited for post-processing  
 FAM ----- AN, PI and PRAI in table, plus Patent Family data  
 FBIB ----- AN, BIB, plus Patent FAM  
 IND ----- Indexing data  
 IPC ----- International Patent Classifications  
 MAX ----- ALL, plus Patent FAM, RE  
 PATS ----- PI, SO  
 SAM ----- CC, SX, TI, ST, IT  
 SCAN ----- CC, SX, TI, ST, IT (random display, no answer numbers;  
           SCAN must be entered on the same line as the DISPLAY,  
           e.g., D SCAN or DISPLAY SCAN)  
 STD ----- BIB, IPC, and NCL  
  
 IABS ----- ABS, indented with text labels  
 IALL ----- ALL, indented with text labels  
 IBIB ----- BIB, indented with text labels  
 IMAX ----- MAX, indented with text labels  
 ISTD ----- STD, indented with text labels  
  
 OBIB ----- AN, plus Bibliographic Data (original)  
 OIBIB ----- OBIB, indented with text labels  
  
 SBIB ----- BIB, no citations  
 SIBIB ----- IBIB, no citations  
  
 HIT ----- Fields containing hit terms  
 HITIND ----- IC, ICA, ICI, NCL, CC and index field (ST and IT)  
               containing hit terms  
 HITRN ----- HIT RN and its text modification  
 HITSTR ----- HIT RN, its text modification, its CA index name, and  
               its structure diagram  
 FHITSTR ----- First HIT RN, its text modification, its CA index name, and  
               its structure diagram  
 KWIC ----- Hit term plus 20 words on either side  
 OCC ----- Number of occurrence of hit term and field in which it occurs

To display a particular field or fields, enter the display field codes. For a list of the display field codes, enter HELP DFIELDS at an arrow prompt (=>). Examples of formats include: TI; TI,AU; BIB,ST; TI,IND; TI,SO. You may specify the format fields in any order and the information will be displayed in the same order as the format specification.

All of the formats (except for SAM, SCAN, HIT, HITIND, HITRN, HITSTR, FHITSTR, KWIC, and OCC) may be used with DISPLAY ACC to view a specified Accession Number.

ENTER DISPLAY FORMAT (BIB):bib

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L38 ANSWER 13 OF 36331 CA COPYRIGHT 2001 ACS
AN 134:220902 CA
TI Activated coagulation/fibrinolysis system and platelet function in acute
      thrombotic stroke patients with increased C-reactive protein
      levels
AU Tohgi, H.; Konno, S.; Takahashi, S.; Koizumi, D.; Kondo, R.; Takahashi,
H.
CS Department of Neurology, Iwate Medical University, Morioka, Iwate,
  
```

027-8505, Japan  
SO Thromb. Res. (2000), 100(5), 373-379  
CODEN: THBRAA; ISSN: 0049-3848  
PB Elsevier Science Inc.  
DT Journal  
LA English  
RE.CNT 32  
RE  
(1) Berk, B; Am J Cardiol 1990, V65, P168 MEDLINE  
(2) Booth, N; Br J Haematol 1988, V70, P327 MEDLINE  
(3) Bova, I; Stroke 1996, V27, P2204 MEDLINE  
(22) Ridker, P; Circulation 1998, V98, P731 CA  
(23) Ridker, P; N Engl J Med 1997, V336, P973 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

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L2 1 S PMSF  
L3 1 S APMSF  
L4 1 S L3  
L5 5 S ANTIPAIN  
L6 170 S ANTITHROMBIN  
L7 11 S LEUPEPTIN  
L8 9 S DICHLOROCOUMARIN  
L9 2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

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L12 24 S L3  
L13 343 S L5  
L14 4326 S L6  
L15 171 S L7  
L16 16 S L8  
L17 372 S L9  
E NEUROPATHY  
L18 5490 S E3  
E EPILEPSY  
L19 10805 S E3-E12  
E SEIZURE  
L20 14562 S E3-E6  
E HYPOXIA  
L21 24801 S E3  
E STROKE  
L22 11880 S E3  
L23 0 S L17 AND L18  
L24 0 S L17 AND L19  
L25 0 S L17 AND L20  
L26 0 S L17 AND L21  
L27 0 S L17 AND L22  
L28 0 S L16 AND L18  
L29 25960 S L18 OR L19 OR L20  
L30 0 S L29 AND L16  
L31 0 S L29 AND L15

L32            4 S L29 AND L14  
L33            0 S L29 AND L13  
L34            0 S L29 AND L12  
L35            30 S L29 AND L11  
L36            0 S L29 AND L10  
L37            350 S L21 AND L22  
L38            36331 S L21 OR L22  
L39            4 S L38 AND L10  
L40            2 S L38 AND L11  
L41            0 S L38 AND L12  
L42            1 S L38 AND L13

=> d 142 1

L42 ANSWER 1 OF 1 CA COPYRIGHT 2001 ACS  
AN 117:103472 CA  
TI A novel cytotoxicity screening assay using a multiwell fluorescence scanner  
AU Nieminen, Anna Liisa; Gores, Gregory J.; Bond, John M.; Imberti, Roberto; Herman, Brian; Lemasters, John J.  
CS Sch. Med., Univ. North Carolina, Chapel Hill, NC, 27599-7090, USA  
SO Toxicol. Appl. Pharmacol. (1992), 115(2), 147-55  
CODEN: TXAPAA; ISSN: 0041-008X  
DT Journal  
LA English

=> s 138 and 114

L43            46 L38 AND L14

=> d 143 20-46

L43 ANSWER 20 OF 46 CA COPYRIGHT 2001 ACS  
AN 130:276486 CA  
TI Antithrombin-III activity and the efficacy of heparin in progressing ischemic **stroke**  
AU Roden-Jullig, Asa; Britton, Mona; Svensson, Jan  
CS Division of Internal Medicine, Karolinska Institutet Danderyd Hospital, Danderyd, S-182 88, Swed.  
SO Clin. Appl. Thromb./Hemostasis (1998), 4(2), 129-132  
CODEN: CATHF9; ISSN: 1076-0296  
PB Lippincott-Raven Publishers  
DT Journal  
LA English  
RE.CNT 24  
RE  
(1) Altes, A; Acta Haematol 1995, V94, P10 MEDLINE  
(3) Britton, M; Stroke 1985, V16, P629 MEDLINE  
(5) Davalos, A; Neurol 1990, V40, P1865 MEDLINE  
(20) Takano, K; Thromb Res 1990, V58, P481 CA  
(23) van Wersch, J; Eur J Clin Chem Clin Biochem 1993, V31, P575 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L43 ANSWER 21 OF 46 CA COPYRIGHT 2001 ACS

AN 130:47481 CA  
TI Preparation of modified, low-molecular-weight heparin that inhibits clot-associated coagulation factors  
IN Weitz, Jeffrey; Hirsh, Jack  
PA Hamilton Civic Hospitals Research Development, Inc., Can.  
SO PCT Int. Appl., 51 pp.

CODEN: PIXXD2

DT Patent  
LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9855515	A1	19981210	WO 1998-CA548	19980605
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	AU 9877538	A1	19981221	AU 1998-77538	19980605
	EP 986581	A1	20000322	EP 1998-925356	19980605
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
PRAI	US 1997-72098		19970606		
	WO 1998-CA548		19980605		

RE.CNT 6

RE

- (1) Bioiberica SA; EP 0337327 A 1989 CA
- (2) Hamilton Civic Hospitals Research Development Inc; EP 0735050 A 1996 CA
- (3) Novo Industri A/S; EP 0244235 A 1987 CA
- (4) Rhone-Poulenc Rorer SA; EP 0511075 A 1992 CA
- (5) Rhone-Poulenc Rorer SA; WO 9316112 A 1993 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L43 ANSWER 22 OF 46 CA COPYRIGHT 2001 ACS

AN 129:329217 CA

TI Relative influence of age and thrombotic history on hemostatic parameters

AU Javorschi, S.; Richard-Harston, S.; Labrouche, S.; Manciet, G.;  
Freyburger, G.

CS Laboratoire d'Hematologie, Hopital Pellegrin, Bordeaux, 33076, Fr.

SO Thromb. Res. (1998), 91(5), 241-248

CODEN: THBRAA; ISSN: 0049-3848

PB Elsevier Science Inc.

DT Journal

LA English

L43 ANSWER 23 OF 46 CA COPYRIGHT 2001 ACS

AN 129:58856 CA

TI Compositions and methods for inhibiting thrombogenesis

IN Weitz, Jeffrey I.; Hirsh, Jack; Young, Edward

PA Hamilton Civic Hospitals Research Development Inc., Can.

SO U.S., 65 pp. Cont.-in-part of U. S. 5,744,457.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 4

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 5763427	A	19980609	US 1996-624327	19960329
	US 5744457	A	19980428	US 1995-540324	19951006
	AU 9651400	A1	19961016	AU 1996-51400	19960329
	JP 11506420	T2	19990608	JP 1996-528734	19960329
	US 6001820	A	19991214	US 1997-870528	19970606
	NO 9704500	A	19971128	NO 1997-4500	19970929
PRAI	US 1995-412332		19950331		
	US 1995-540324		19951006		
	US 1996-624327		19960329		

WO 1996-CA190 19960329  
OS MARPAT 129:58856

L43 ANSWER 24 OF 46 CA COPYRIGHT 2001 ACS  
AN 129:26531 CA  
TI Analysis of lipoprotein(a) and coagulation-fibrinolysis system in the stroke types or recurrences of the cerebral thrombosis  
AU Kashiwaya, Mitsuru; Konno, Shu; Takahashi, Hiroaki  
CS Sch. Med., Iwate Med. Univ., Morioka, 020-8505, Japan  
SO Iwate Igaku Zasshi (1998), 50(1), 65-71  
CODEN: IIZAAX; ISSN: 0021-3284  
PB Iwate Igakkai  
DT Journal  
LA Japanese

L43 ANSWER 25 OF 46 CA COPYRIGHT 2001 ACS  
AN 129:3340 CA  
TI Pathophysiology and clinical management of fetus with intrauterine growth retardation  
AU Takeda, Yoshihiko  
CS Dep. Gynecol., Tokyo Women's Med. Coll., Tokyo, 162-8666, Japan  
SO Tokyo Joshi Ika Daigaku Zasshi (1998), 68(4), 161-170  
CODEN: TJIZAF; ISSN: 0040-9022  
PB Tokyo Joshi Ika Daigaku Gakkai  
DT Journal; General Review  
LA Japanese

L43 ANSWER 26 OF 46 CA COPYRIGHT 2001 ACS  
AN 128:312895 CA  
TI Compositions and methods for inhibiting thrombogenesis  
IN Weitz, Jeffrey I.; Hirsh, Jack; Young, Edward  
PA Hamilton Civic Hospitals Research Development Inc., Can.  
SO U.S., 67 pp. Cont.-in-part of U.S. Ser. No. 412,332, abandoned.  
CODEN: USXXAM  
DT Patent  
LA English  
FAN.CNT 4

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 5744457	A	19980428	US 1995-540324	19951006
	WO 9629973	A2	19961003	WO 1996-CA190	19960329
	WO 9629973	A3	19961219		
	W:	AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI			
	RW:	KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML			
	CA 2217054	AA	19961003	CA 1996-2217054	19960329
	AU 9651400	A1	19961016	AU 1996-51400	19960329
	US 5763427	A	19980609	US 1996-624327	19960329
	CN 1186502	A	19980701	CN 1996-194354	19960329
	JP 11506420	T2	19990608	JP 1996-528734	19960329
	EP 735050	A2	19961002	EP 1996-302311	19960401
	EP 735050	A3	19970122		
	R:	AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE			
	GB 2299998	A1	19961023	GB 1996-6881	19960401
	GB 2299998	B2	19970326		
	US 6001820	A	19991214	US 1997-870528	19970606
	NO 9704500	A	19971128	NO 1997-4500	19970929
PRAI	US 1995-412332	19950331			
	US 1995-485872	19950607			

US 1995-540324 19951006  
US 1996-624327 19960329  
WO 1996-CA190 19960329  
OS MARPAT 128:312895

L43 ANSWER 27 OF 46 CA COPYRIGHT 2001 ACS  
AN 128:305295 CA  
TI Histidine-proline-rich glycoprotein as a plasma pH sensor. Modulation of its interaction with glycosaminoglycans by pH and metals  
AU Borza, Dorin-Bogdan; Morgan, William T.  
CS Division of Molecular Biology and Biochemistry, School of Biological Sciences, University of Missouri, Kansas City, MO, 64110, USA  
SO J. Biol. Chem. (1998), 273(10), 5493-5499  
CODEN: JBCHA3; ISSN: 0021-9258  
PB American Society for Biochemistry and Molecular Biology  
DT Journal  
LA English

L43 ANSWER 28 OF 46 CA COPYRIGHT 2001 ACS  
AN 128:97817 CA  
TI Effects of oral and transdermal estrogen/progesterone regimens on blood coagulation and fibrinolysis in postmenopausal women: a randomized controlled trial  
AU Scarabin, Pierre-Yves; Alhenc-Gelas, Martine; Plu-Bureau, Genevieve; Taisne, Pascale; Agher, Rachid; Aiach, Martine  
CS INSERM-Cardiovascular Epidemiology Unit U258, Hopital Broussais, Paris, Fr.  
SO Arterioscler., Thromb., Vasc. Biol. (1997), 17(11), 3071-3078  
CODEN: ATVBFA; ISSN: 1079-5642  
PB American Heart Association  
DT Journal  
LA English

L43 ANSWER 29 OF 46 CA COPYRIGHT 2001 ACS  
AN 125:325404 CA  
TI Hemostatic studies in carbohydrate-deficient glycoprotein syndrome type I  
AU Fiumara, A.; Barone, R.; Buttitta, P.; Musso, R.; Pavone, L.; Nigro, F.; Jaeken, J.  
CS Department Pediatrics, University Catania, Catania, Italy  
SO Thromb. Haemostasis (1996), 76(4), 502-504  
CODEN: THHADQ; ISSN: 0340-6245  
DT Journal  
LA English

L43 ANSWER 30 OF 46 CA COPYRIGHT 2001 ACS  
AN 125:317356 CA  
TI Heparin preparations for inhibiting thrombogenesis  
IN Weitz, Jeffrey I.; Hirsh, Jack; Young, Edward  
PA Hamilton Civic Hospitals Research Development, Inc., Can.  
SO Eur. Pat. Appl., 76 pp.  
CODEN: EPXXDW  
DT Patent  
LA English

FAN.CNT 4

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 735050	A2	19961002	EP 1996-302311	19960401
	EP 735050	A3	19970122		
	R: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
	US 5744457	A	19980428	US 1995-540324	19951006
	AU 9651400	A1	19961016	AU 1996-51400	19960329
	JP 11506420	T2	19990608	JP 1996-528734	19960329

NO 9704500 A 19971128 NO 1997-4500 19970929  
PRAI US 1995-412332 19950331  
US 1995-485872 19950607  
US 1995-540324 19951006  
WO 1996-CA190 19960329

L43 ANSWER 31 OF 46 CA COPYRIGHT 2001 ACS  
AN 125:104648 CA  
TI Antithrombin III prevents blood pressure elevation and proteinuria induced by high salt intake in pregnant stroke-prone spontaneously hypertensive rats  
AU Shinya, Hiroshi; Yamanaga, Katsumi; Akira, Toshiaki; Uchida, Takeshi; Yaguchi, Masafumi; Watanabe, Masahiro; Kagitani, Yoshio  
CS Pharmacology Labs., Green Cross Corp., Osaka, 573, Japan  
SO Biol. Pharm. Bull. (1996), 19(6), 819-823  
CODEN: BPBLEO; ISSN: 0918-6158  
DT Journal  
LA English

L43 ANSWER 32 OF 46 CA COPYRIGHT 2001 ACS  
AN 125:83683 CA  
TI Complex functional and structural coagulation abnormalities in the carbohydrate-deficient glycoprotein syndrome type I  
AU Stibler, H.; Holzbach, U.; Tengborn, L.; Kristiansson, B.  
CS Department Neurology, Karolinska Hospital, Stockholm, S-171 76, Swed.  
SO Blood Coagulation Fibrinolysis (1996), 7(2), 118-126  
CODEN: BLFIE7; ISSN: 0957-5235  
DT Journal  
LA English

L43 ANSWER 33 OF 46 CA COPYRIGHT 2001 ACS  
AN 125:80523 CA  
TI .alpha.1-antitrypsin variants carrying thrombin-specificity peptides from antithrombin III that are inactive against activated protein C  
IN Hopkins, Paul C. R.; Carrell, Robin; Crowther, Damian; Stone, Stuart  
PA Ppl Therapeutics (Scotland) Ltd., UK  
SO PCT Int. Appl., 50 pp.  
CODEN: PIXXD2  
DT Patent  
LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9610638	A1	19960411	WO 1995-GB2155	19950912
	W:	AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LU, LV, MD, MG, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TT			
	RW:	KE, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG			
	AU 9534794	A1	19960426	AU 1995-34794	19950912
	ZA 9507852	A	19970318	ZA 1995-7852	19950918
PRAI	GB 1994-19804		19940930		
	GB 1995-2138		19950203		
	WO 1995-GB2155		19950912		

L43 ANSWER 34 OF 46 CA COPYRIGHT 2001 ACS  
AN 124:6115 CA  
TI Lipoprotein(a) and hemostasis activation markers in angina pectoris  
AU Tsakiris, D. A.; Riesen, W. F.; Marbet, G. A.  
CS Dep. Zentrallab., Univ. Basel, Basel, CH-4031, Switz.

SO Dtsch. Med. Wochenschr. (1995), 120(33), 1109-13  
CODEN: DMWOAX; ISSN: 0012-0472  
DT Journal  
LA English

L43 ANSWER 35 OF 46 CA COPYRIGHT 2001 ACS  
AN 121:149109 CA  
TI Treatment of neurodegenerative diseases with thrombin inhibitors  
IN Friedrich, Thomas  
PA BASF A.-G., Germany  
SO Ger. Offen., 4 pp.  
CODEN: GWXXBX  
DT Patent  
LA German  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 4303646	A1	19940811	DE 1993-4303646	19930209
	WO 9417821	A1	19940818	WO 1994-EP259	19940129
	W: CA, JP, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	CA 2153420	AA	19940818	CA 1994-2153420	19940129
	EP 683674	A1	19951129	EP 1994-906174	19940129
	R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, NL, PT, SE				
	JP 08507047	T2	19960730	JP 1994-517604	19940129
PRAI	DE 1993-4303646		19930209		
	WO 1994-EP259		19940129		

L43 ANSWER 36 OF 46 CA COPYRIGHT 2001 ACS  
AN 120:51584 CA  
TI Changes of von Willebrand factor and antithrombin III levels in acute stroke: Difference between thrombotic and hemorrhagic stroke  
AU Liu, Longbin; Lin, Zhusan; Shen, Zeshuang  
CS 2nd Affil. Hosp., Hunan Med. Univ., Peop. Rep. China  
SO Thromb. Res. (1993), 72(4), 353-8  
CODEN: THBRAA; ISSN: 0049-3848  
DT Journal  
LA English

L43 ANSWER 37 OF 46 CA COPYRIGHT 2001 ACS  
AN 118:94067 CA  
TI Effect of long-term mesoglycan treatment on fibrinogen plasma levels in patients with ischemic cerebrovascular disease  
AU Orefice, G.; Troisi, E.; Selvaggio, M.; Vecchione, V.; Rubino, S.; Provitera, V.; Carrieri, P. B.  
CS 2nd Med. Sch., Univ. Naples, Naples, Italy  
SO Curr. Ther. Res. (1992), 52(5), 666-70  
CODEN: CTCEA9; ISSN: 0011-393X  
DT Journal  
LA English

L43 ANSWER 38 OF 46 CA COPYRIGHT 2001 ACS  
AN 118:78363 CA  
TI Effects of hypoxia on heparan sulfate in bovine aortic and pulmonary artery endothelial cells  
AU Karlinsky, Joel B.; Rounds, Sharon; Farber, Harrison W.  
CS Sch. Med., Boston Univ., Boston, MA, USA  
SO Circ. Res. (1992), 71(4), 782-9  
CODEN: CIRUAL; ISSN: 0009-7330  
DT Journal  
LA English

L43 ANSWER 39 OF 46 CA COPYRIGHT 2001 ACS  
AN 118:16315 CA  
TI Chimeric molecule with plasminogen activator activity and affinity for  
atherosclerotic plaques  
IN Loscalzo, Joseph; Pasche, Boris  
PA Brigham and Women's Hospital, USA  
SO PCT Int. Appl., 29 pp.  
CODEN: PIXXD2  
DT Patent  
LA English  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9218139	A1	19921029	WO 1992-US3009	19920409
	W: AU, CA, JP			RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, MC, NL, SE	
	AU 9218730	A1	19921117	AU 1992-18730	19920409
PRAI	US 1991-682070	19910409			
	WO 1992-US3009	19920409			

L43 ANSWER 40 OF 46 CA COPYRIGHT 2001 ACS  
AN 117:168788 CA  
TI Protein S deficiency in middle-aged women with **stroke**  
AU Green, David; Otoya, Jorge; Oriba, Howard; Rovner, Richard  
CS Med. Sch., Northwestern Univ., Chicago, IL, 60611, USA  
SO Neurology (1992), 42(5), 1029-33  
CODEN: NEURAI; ISSN: 0028-3878  
DT Journal  
LA English

L43 ANSWER 41 OF 46 CA COPYRIGHT 2001 ACS  
AN 114:226557 CA  
TI Circadian variations of platelet aggregability and fibrinolytic activity  
in healthy subjects  
AU Jovicic, A.; Mandic, S.  
CS Clin. Neurol., Mil. Med. Acad., Belgrade, Yugoslavia  
SO Thromb. Res. (1991), 62(1-2), 65-74  
CODEN: THBRAA; ISSN: 0049-3848  
DT Journal  
LA English

L43 ANSWER 42 OF 46 CA COPYRIGHT 2001 ACS  
AN 113:75820 CA  
TI Hypercoagulability in acute ischemic **stroke**: analysis of the  
extrinsic coagulation reactions in plasma by a highly sensitive automated  
method  
AU Takano, Kentaro; Yamaguchi, Takenori; Okada, Yasushi; Uchida, Kagehiro;  
Kisiel, Walter; Kato, Hisao  
CS Res. Inst., Natl. Cardiovasc. Cent., Osaka, Japan  
SO Thromb. Res. (1990), 58(5), 481-91  
CODEN: THBRAA; ISSN: 0049-3848  
DT Journal  
LA English

L43 ANSWER 43 OF 46 CA COPYRIGHT 2001 ACS  
AN 108:34244 CA  
TI Heparin cofactor II: a simple assay method and results of its clinical  
application  
AU Vinazzer, H.; Pangraz, U.  
CS Blood Coagulation Lab., Linz, Austria  
SO Thromb. Res. (1987), 48(2), 153-60  
CODEN: THBRAA; ISSN: 0049-3848  
DT Journal

LA English

L43 ANSWER 44 OF 46 CA COPYRIGHT 2001 ACS  
AN 89:3808 CA  
TI Some characteristics of blood coagulation in people under high-altitude conditions determined by the processing of data in a computer  
AU Isabaeva, V. A.; Prizhivoit, G. N.; Gurovich, T. Ts.; Prizhivoit, Ya. I.; Shablovskii, V. I.  
CS Kirg. Med. Inst., Frunze, USSR  
SO Tr. Kirg. Gos. Med. Inst. (1976), 110, 87-94  
CODEN: TKRMAS; ISSN: 0371-8778  
DT Journal  
LA Russian

L43 ANSWER 45 OF 46 CA COPYRIGHT 2001 ACS  
AN 89:3807 CA  
TI The blood clotting system during high-altitude adaptation and readaptation  
AU Isabaeva, V. A.; Ponomareva, T. A.  
CS Kirg. Med. Inst., Frunze, USSR  
SO Tr. Kirg. Gos. Med. Inst. (1976), 110, 77-87  
CODEN: TKRMAS; ISSN: 0371-8778  
DT Journal  
LA Russian

L43 ANSWER 46 OF 46 CA COPYRIGHT 2001 ACS  
AN 85:91728 CA  
TI Blood coagulation and plasma fibrinolytic enzyme system pathophysiology in **stroke**  
AU Fletcher, Anthony P.; Alkjaersig, Norma; Davies, Andrew; Lewis, Martin; Brooks, John; Hardin, William; Landau, William; Raichle, Marcus E.  
CS Sch. Med., Washington Univ., St. Louis, Mo., USA  
SO Stroke (1976), 7(4), 337-48  
CODEN: SJCCA7  
DT Journal  
LA English

=> d 143 46 42 36 30 26 24 all

L43 ANSWER 46 OF 46 CA COPYRIGHT 2001 ACS  
AN 85:91728 CA  
TI Blood coagulation and plasma fibrinolytic enzyme system pathophysiology in **stroke**  
AU Fletcher, Anthony P.; Alkjaersig, Norma; Davies, Andrew; Lewis, Martin; Brooks, John; Hardin, William; Landau, William; Raichle, Marcus E.  
CS Sch. Med., Washington Univ., St. Louis, Mo., USA  
SO Stroke (1976), 7(4), 337-48  
CODEN: SJCCA7  
DT Journal  
LA English  
CC 14-7 (Mammalian Pathological Biochemistry)  
AB Plasma fibrinogen chromatog. is a method for quantification of high mol. wt. fibrinogen complexes (HMWFC), native fibrinogen and other fibrinogen derivs. in plasma. The method distinguishes between subjects with normal and pathol. rates of fibrin formation. Serial std. blood coagulation assays, including plasma fibrinogen chromatog., and neurol. studies were performed on 220 patients admitted to a **stroke** unit. Findings from patients with cerebral infarction were compared against those of 3 control groups: normals, a **stroke** control group, and a

**stroke** risk factor group. Plasma HMWFC findings were higher in the **stroke** risk factor group than in the normals. Plasma HMWFC values were higher in the cerebral infarction patients than in any of the control groups, and plasma fibrinogen, plasminogen, .alpha.1-antitrypsin and .alpha.2-macroglobulin also were higher in the patients. The greater the degree of initial neurrol. deficit, the greater were plasma HMWFC values, and high HMWFC values were associated with poor clin. outcome. Plasma HMWFC values were higher in patients with intracerebral hemorrhage,

subarachnoid hemorrhage, and cerebral embolism. Thus, a high proportion of **stroke** patients have coagulopathy, characterized by pathol. enhancement of fibrin formation.

ST **stroke** plasma fibrinogen chromatog; blood coagulation chromatog

**stroke**

IT Macroglobulins

RL: BIOL (Biological study)  
(.alpha.2-, in brain circulatory disease)

IT Brain, disease or disorder  
(circulatory, blood coagulation and fibrinolysis in)

IT Fibrinolysis

Fibrinogens  
RL: BIOL (Biological study)  
(in brain circulatory disease)

IT 9000-94-6

RL: BIOL (Biological study)  
(III, in brain circulatory disease)

IT 9001-91-6 9041-92-3

RL: BIOL (Biological study)  
(in brain circulatory disease)

L43 ANSWER 42 OF 46 CA COPYRIGHT 2001 ACS

AN 113:75820 CA

TI Hypercoagulability in acute ischemic **stroke**: analysis of the extrinsic coagulation reactions in plasma by a highly sensitive automated method

AU Takano, Kentaro; Yamaguchi, Takenori; Okada, Yasushi; Uchida, Kagehiro; Kisiel, Walter; Kato, Hisao

CS Res. Inst., Natl. Cardiovasc. Cent., Osaka, Japan

SO Thromb. Res. (1990), 58(5), 481-91

CODEN: THBRAA; ISSN: 0049-3848

DT Journal

LA English

CC 14-6 (Mammalian Pathological Biochemistry)

AB The coagulability of plasma from 63 patients with acute ischemic **stroke** (cerebral thrombosis and cerebral embolism) was analyzed by an automated method for prothrombin time using a fluorogenic peptide substrate. The fluorogenic prothrombin time (FPT) of plasma collected within 48 h after onset, as expressed as percent of control plasma, was significantly higher in cerebral thrombosis than in an age-matched control

group. The high values of FPT in cerebral thrombosis patients were obsd. until the 30th day after onset. On the other hand, FPT values in cerebral

embolism patients were not significantly different than that of the control group. Factor VII activity levels in cerebral thrombosis patients

were significantly higher than those of the control group and cerebral embolism patients, while levels of factor X activity were not significantly different among these groups. Although FPT and factor VII activity in these **stroke** patients did not significantly correlate, factor VII activity did correlate well with factor VII antigen.

Decreased levels of antithrombin III and elevated levels of FDP and

.alpha.2-antiplasmin-plasmin complexes were obsd. only in cerebral embolism patients. The findings strongly suggest that patients with cerebral thrombosis have been in a hypercoagulable state before the onset of symptoms, which was caused in part by an increase of factor VII activity/antigen, and in part by other unknown mechanisms. In contrast, patients with cerebral embolism were in a low grade consumptive coagulopathy.

- ST brain thrombosis blood coagulation factor VII; hypercoagulability  
stroke blood coagulation factor VII
- IT Fibrinogen degradation products  
RL: BIOL (Biological study)  
(in ischemic stroke from cerebral embolism, of humans, low grade consumptive coagulopathy in relation to)
- IT Embolism  
(ischemic stroke of brain from, antithrombin III and fibrinogen degrdn. products and .alpha.2-antiplasmin-plasmin complexes of blood plasma and low grade consumptive coagulopathy in relation to, of humans)
- IT Thrombosis  
(ischemic stroke of brain from, blood-coagulation factor VII of blood plasma and hypercoagulability in relation to, of human)
- IT Blood coagulation  
(disorder, disseminated intravascular, antithrombin III and fibrinogen degrdn. product and .alpha.2-antiplasmin-plasmin complexes of blood plasma in ischemic stroke from cerebral embolism of human in relation to)
- IT Blood coagulation  
(disorder, hypercoagulability, blood-coagulation factor VII of blood plasma in ischemic stroke from thrombosis of humans in relation to)
- IT Brain, disease or disorder  
(embolism, ischemic stroke from, antithrombin III and fibrinogen degrdn. products and .alpha.2-antiplasmin-plasmin complexes of blood plasma and low grade consumptive coagulopathy in relation to, of humans)
- IT Brain, disease or disorder  
(stroke, from embolism or thrombosis, extrinsic coagulation factors of blood plasma in, of humans, coagulopathy in relation to)
- IT Brain, disease or disorder  
(thrombosis, ischemic stroke from, blood-coagulation factor VII and hypercoagulability in relation to, of humans)
- IT 9000-94-6, Antithrombin  
RL: BIOL (Biological study)  
(in ischemic stroke from cerebral embolism, of humans, low grade consumptive coagulopathy in relation to)
- IT 9001-25-6, Blood-coagulation factor VII  
RL: BIOL (Biological study)  
(in ischemic stroke from cerebral thrombosis, of humans, hypercoagulability in relation to)
- IT 9001-90-5, Plasmin  
RL: BIOL (Biological study)  
(.alpha.2-antiplasmin complexes, in ischemic stroke from cerebral embolism, of humans, low grade consumptive coagulopathy in relation to)

L43 ANSWER 36 OF 46 CA COPYRIGHT 2001 ACS

AN 120:51584 CA

TI Changes of von Willebrand factor and antithrombin III levels in acute stroke: Difference between thrombotic and hemorrhagic stroke

AU Liu, Longbin; Lin, Zhusan; Shen, Zeshuang

CS 2nd Affil. Hosp., Hunan Med. Univ., Peop. Rep. China

SO Thromb. Res. (1993), 72(4), 353-8

DT Journal  
 LA English  
 CC 14-6 (Mammalian Pathological Biochemistry)  
 AB In the present study, the authors measured plasma von Willebrand factor (vWF) levels, concns. and activities. of Antithrombin III (ATIII) in the acute phase of thrombotic and hemorrhagic **stroke** prior to any therapy. The authors' results demonstrate that vWF levels are increased in both thrombotic and hemorrhagic **stroke**, and that vWF and ATIII levels differ between thrombotic and hemorrhagic **stroke** in patients with high incidence of atherosclerosis.  
 ST von Willebrand antithrombin thrombotic hemorrhagic **stroke**  
 IT Brain, disease  
     (hemorrhagic **stroke**, antithrombin III and von Willebrand factor of blood plasma in, in human)  
 IT Brain, disease  
     (thrombotic **stroke**, antithrombin III and von Willebrand factor of blood plasma in, in human)  
 IT 109319-16-6  
     RL: BIOL (Biological study)  
       (of blood plasma, in human hemorrhagic and thrombotic **stroke**, antithrombin III in relation to)  
 IT 9000-94-6, Antithrombin III  
     RL: BIOL (Biological study)  
       (of blood plasma, in human hemorrhagic and thrombotic **stroke**, von Willebrand factor in relation to)

L43 ANSWER 30 OF 46 CA COPYRIGHT 2001 ACS  
 AN 125:317356 CA  
 TI Heparin preparations for inhibiting thrombogenesis  
 IN Weitz, Jeffrey I.; Hirsh, Jack; Young, Edward  
 PA Hamilton Civic Hospitals Research Development, Inc., Can.  
 SO Eur. Pat. Appl., 76 pp.  
 CODEN: EPXXDW  
 DT Patent  
 LA English  
 IC ICM C08B037-10  
 ICS A61K031-725  
 CC 1-8 (Pharmacology)  
 Section cross-reference(s): 33  
 FAN.CNT 4

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 735050	A2	19961002	EP 1996-302311	19960401
	EP 735050	A3	19970122		
	R: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
	US 5744457	A	19980428	US 1995-540324	19951006
	AU 9651400	A1	19961016	AU 1996-51400	19960329
	JP 11506420	T2	19990608	JP 1996-528734	19960329
	NO 9704500	A	19971128	NO 1997-4500	19970929
PRAI	US 1995-412332	19950331			
	US 1995-485872	19950607			
	US 1995-540324	19951006			
	WO 1996-CA190	19960329			
AB	Comps. are provided for inactivating thrombin bound to fibrin within a thrombus or clot, whereby the ability of clot-bound thrombin to catalytically promote further clot accretion is a substantially diminished				
	or eliminated. The compns. are particularly useful for preventing thrombosis in the circuit of cardiac bypass app. and in patients undergoing renal dialysis, and for treating patients suffering from or at risk of suffering from thrombus-related cardiovascular conditions, such as				

unstable angina, acute myocardial infarction (heart attack), cerebrovascular accidents (**stroke**), pulmonary embolism, deep vein thrombosis, arterial thrombosis, etc. The compns. comprise agents which activate heparin cofactor II-mediated inhibition of thrombin and having minimal affinity for antithrombin III. Preferred agents are low mol. wt. heparin prepns. (MW of 3,000-8,000) prep'd. by depolymerg. heparin using nitrous acid, oxidizing the resultant product with periodate and reducing it with borohydride. The product has its non-sulfated uronic acid residues in open ring form and it substantially free of aldehyde groups.

- ST heparin deriv prep'n thrombogenesis inhibition  
IT Blood platelet  
    (factor Xa binding to; heparin prepns. for inhibiting thrombogenesis)  
IT Anticoagulants and Antithrombotics  
Cardiovascular agents  
Drug interactions  
Oxidizing agents  
Pharmaceutical dosage forms  
Reducing agents  
    .heparin prepns. for inhibiting thrombogenesis)  
IT Desmins  
RL: BAC (Biological activity or effector, except adverse); THU  
(Therapeutic use); BIOL (Biological study); USES (Uses)  
    .heparin prepns. for inhibiting thrombogenesis)  
IT Depolymerization  
    (nitrous acid; heparin prepns. for inhibiting thrombogenesis)  
IT Anhydrides  
RL: RCT (Reactant)  
    (oxidizing agent; heparin prepns. for inhibiting thrombogenesis)  
IT Uronic acids  
RL: PRP (Properties)  
    (polyanionic carbohydrate contg.; heparin prepns. for inhibiting thrombogenesis)  
IT Fibrins  
RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
    (poymn.; heparin prepns. for inhibiting thrombogenesis)  
IT Hydrides  
RL: RCT (Reactant)  
    (reducing agent; heparin prepns. for inhibiting thrombogenesis)  
IT Thrombus and Blood clot  
    (thrombin bound to; heparin prepns. for inhibiting thrombogenesis)  
IT Carbohydrates and Sugars, biological studies  
RL: BAC (Biological activity or effector, except adverse); THU  
(Therapeutic use); BIOL (Biological study); USES (Uses)  
    (anionic, heparin prepns. for inhibiting thrombogenesis)  
IT Circulation  
    (extracorporeal, cardiopulmonary bypass, heparin prepns. for  
inhibiting  
    thrombogenesis)  
IT 9002-05-5, Blood coagulation factor Xa  
RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
    (binding to platelet surface; heparin prepns. for inhibiting  
    thrombogenesis)  
IT 9002-04-4, Thrombin  
RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
    (clot-bound; heparin prepns. for inhibiting thrombogenesis)  
IT 7782-77-6, Nitrous acid  
RL: RCT (Reactant)  
    (depolym.; heparin prepns. for inhibiting thrombogenesis)  
IT 8001-27-2, Hirudin  
RL: ADV (Adverse effect, including toxicity); BAC (Biological activity or  
effector, except adverse); THU (Therapeutic use); BIOL (Biological  
study);

USES (Uses)  
 (heparin prepns. for inhibiting thrombogenesis)

IT 9005-49-6, Heparin, biological studies 9005-49-6D, Heparin, derivs.  
 RL: BAC (Biological activity or effector, except adverse); THU  
 (Therapeutic use); BIOL (Biological study); USES (Uses)  
 (heparin prepns. for inhibiting thrombogenesis)

IT 9000-94-6, Antithrombin III 37203-61-5, Blood coagulation factor  
 XIA 37316-87-3, Blood coagulation factor IXa 81604-65-1, Heparin  
 cofactor II  
 RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
 (heparin prepns. for inhibiting thrombogenesis)

IT 50-81-7, Ascorbic acid, reactions 67-68-5, Dimethyl sulfoxide,  
 reactions  
 546-67-8, Lead tetraacetate 7790-28-5, Sodium periodate  
 RL: RCT (Reactant)  
 (oxidizing agent; heparin prepns. for inhibiting thrombogenesis)

IT 41107-82-8  
 RL: PRP (Properties)  
 (polyanionic carbohydrate contg.; heparin prepns. for inhibiting  
 thrombogenesis)

IT 302-01-2, Hydrazine, reactions 16853-85-3, Lithium aluminum hydride  
 16940-66-2, Sodium borohydride  
 RL: RCT (Reactant)  
 (reducing agent; heparin prepns. for inhibiting thrombogenesis)

L43 ANSWER 26 OF 46 CA COPYRIGHT 2001 ACS

AN 128:312895 CA

TI Compositions and methods for inhibiting thrombogenesis

IN Weitz, Jeffrey I.; Hirsh, Jack; Young, Edward

PA Hamilton Civic Hospitals Research Development Inc., Can.

SO U.S., 67 pp. Cont.-in-part of U.S. Ser. No. 412,332, abandoned.

CODEN: USXXAM

DT Patent

LA English

IC ICM A61K031-725

ICS C08B037-10

NCL 514056000

CC 63-3 (Pharmaceuticals)

Section cross-reference(s): 1

FAN.CNT 4

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 5744457	A	19980428	US 1995-540324	19951006
	WO 9629973	A2	19961003	WO 1996-CA190	19960329
	WO 9629973	A3	19961219		
	W:	AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI			
	RW:	KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML			
	CA 2217054	AA	19961003	CA 1996-2217054	19960329
	AU 9651400	A1	19961016	AU 1996-51400	19960329
	US 5763427	A	19980609	US 1996-624327	19960329
	CN 1186502	A	19980701	CN 1996-194354	19960329
	JP 11506420	T2	19990608	JP 1996-528734	19960329
	EP 735050	A2	19961002	EP 1996-302311	19960401
	EP 735050	A3	19970122		
	R:	AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE			
	GB 2299998	A1	19961023	GB 1996-6881	19960401
	GB 2299998	B2	19970326		
	US 6001820	A	19991214	US 1997-870528	19970606

	NO 9704500	A 19971128	NO 1997-4500	19970929
PRAI	US 1995-412332	19950331		
	US 1995-485872	19950607		
	US 1995-540324	19951006		
	US 1996-624327	19960329		
	WO 1996-CA190	19960329		
OS	MARPAT 128:312895			
AB	The present invention provides compns. and methods for inactivating thrombin bound to fibrin within a thrombus or clot, whereby the ability of			
of	clot-bound thrombin to catalytically promote further clot accretion is substantially diminished or eliminated. The compns. contg. heparin cofactor II-specific catalysts are particularly useful for preventing thrombosis in the circuit of cardiac bypass app. and in patients undergoing renal dialysis, and for treating patients suffering from or at risk of suffering from thrombus-related cardiovascular conditions, such as			
as	unstable angina, acute myocardial infarction (heart attack), cerebrovascular accidents ( <b>stroke</b> ), pulmonary embolism, deep vein thrombosis, arterial thrombosis, etc. The heparin preps. consist of			
ST	the lowest 1/3 mol. wt. fraction isolated from unfractionated heparin.			
IT	antithrombogenic heparin prepn			
IT	Extracorporeal circulation (cardiopulmonary bypass; heparin fractions for inhibiting thrombogenesis)			
IT	Anticoagulants			
	Dialysis			
	Oxidizing agents			
	Reducing agents			
	Thrombolytics (heparin fractions for inhibiting thrombogenesis)			
IT	Anhydrides RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process) (oxidizing agents; heparin fractions for inhibiting thrombogenesis)			
IT	Hydrides RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process) (reducing agents; heparin fractions for inhibiting thrombogenesis)			
IT	81604-65-1, Heparin cofactor II RL: BSU (Biological study, unclassified); BIOL (Biological study) (catalysts for; heparin fractions for inhibiting thrombogenesis)			
IT	9005-49-6, Heparin, biological studies RL: BAC (Biological activity or effector, except adverse); PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses) (heparin fractions for inhibiting thrombogenesis)			
IT	9000-94-6, Antithrombin III RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (heparin fractions for inhibiting thrombogenesis)			
IT	50-81-7, Ascorbic acid, reactions 67-68-5, Dmso, reactions 546-67-8, Lead tetraacetate 7790-28-5, Sodium periodate RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process) (oxidizing agent; heparin fractions for inhibiting thrombogenesis)			
IT	302-01-2, Hydrazine, reactions 16853-85-3, Lithium aluminum hydride 16940-66-2, Sodium borohydride RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process) (reducing agent; heparin fractions for inhibiting thrombogenesis)			

AN 129:26531 CA  
TI Analysis of lipoprotein(a) and coagulation-fibrinolysis system in the stroke types or recurrences of the cerebral thrombosis  
AU Kashiwaya, Mitsuru; Konno, Shu; Takahashi, Hiroaki  
CS Sch. Med., Iwate Med. Univ., Morioka, 020-8505, Japan  
SO Iwate Igaku Zasshi (1998), 50(1), 65-71  
CODEN: IIZAAX; ISSN: 0021-3284  
PB Iwate Igakkai  
DT Journal  
LA Japanese  
CC 14-5 (Mammalian Pathological Biochemistry)  
AB We studied the serum level of lipoprotein(a) and fibrinolysis system in 235 non-embolic cerebral thrombosis patients diagnosed by CT scans at the initial stroke. One hundred and 56 of them were followed up for 3.1 +-. 2.7 yr, and the rate of symptomatic and asymptomatic reinfarcts was also studied. Sixty out of 235 patients (26%) had the Lp(a) levels  $\geq 20$  mg/dL. There was no significant difference in the location of cerebral infarction between patients with Lp(a) levels  $< 20$  mg/dL and those with Lp(a) levels  $\geq 20$  mg/dL. The relative risk of reinfarct in patients with Lp(a)  $\geq 20$  mg/dL compared with those with Lp(a)  $< 20$  mg/dL was significantly greater for symptomatic reinfarct ( $p < 0.05$ ), but not for asymptomatic reinfarct. The concn. of fibrinogen and the thrombin-antithrombin complex were higher in patients with Lp(a)  $\geq 20$  mg/dL than those with Lp(a)  $< 20$  mg/dL in the acute and chronic phases ( $p < 0.05$ ). The plasmin-alpha.2 inhibitor complex and D dimer levels were significantly higher in patients with high-Lp(a) levels in chronic phase ( $p < 0.05$ ). These results suggest that elevated Lp(a) levels predisposed symptomatic reinfarct, and were assocd. with a greater activation of the coagulation system in the acute and chronic phases, and of the fibrinolysis system in the chronic phase in cerebral thrombosis.  
ST lipoprotein a cerebral thrombosis coagulation fibrinolysis; plasmin alpha2  
fibrinogen thrombin antithrombin Lpa  
IT High-density lipoproteins  
RL: BOC (Biological occurrence); BIOL (Biological study); OCCU (Occurrence)  
(cholesterol; serum lipoprotein (a) and coagulation-fibrinolysis system  
in stroke types or recurrences of cerebral thrombosis)  
IT Cerebral artery  
Fibrinolysis  
Stroke  
(serum lipoprotein (a) and coagulation-fibrinolysis system in stroke types or recurrences of cerebral thrombosis)  
IT Blood cholesterol  
Blood triglycerides  
D-dimer (fibrinogen degradation product)  
Fibrinogens  
Lipoprotein(a)  
RL: BOC (Biological occurrence); BIOL (Biological study); OCCU (Occurrence)  
(serum lipoprotein (a) and coagulation-fibrinolysis system in stroke types or recurrences of cerebral thrombosis)  
IT 57-88-5, Cholesterol, biological studies 9000-94-6D,  
Antithrombin III, thrombin complex 9001-90-5D, Plasmin,  
.alpha.2-plasmin  
inhibitor complexes 9002-04-4D, Thrombin, antithrombin III complex  
138757-15-0D, .alpha.2-Plasmin inhibitor, plasmin complexes  
RL: BOC (Biological occurrence); BIOL (Biological study); OCCU (Occurrence)  
(serum lipoprotein (a) and coagulation-fibrinolysis system in

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1 11 S NEUROSERPIN  
L2 1 S PMSF  
L3 1 S APMSF  
L4 1 S L3  
L5 5 S ANTIPAIN  
L6 170 S ANTITHROMBIN  
L7 11 S LEUPEPTIN  
L8 9 S DICHLOROCOUMARIN  
L9 2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

L10 17 S L1  
E NEUROSERPIN  
L11 728 S L2  
L12 24 S L3  
L13 343 S L5  
L14 4326 S L6  
L15 171 S L7  
L16 16 S L8  
L17 372 S L9  
E NEUROPATHY  
L18 5490 S E3  
E EPILEPSY  
L19 10805 S E3-E12  
E SEIZURE  
L20 14562 S E3-E6  
E HYPOXIA  
L21 24801 S E3  
E STROKE  
L22 11880 S E3  
0 S L17 AND L18  
L24 0 S L17 AND L19  
L25 0 S L17 AND L20  
L26 0 S L17 AND L21  
L27 0 S L17 AND L22  
L28 0 S L16 AND L18  
L29 25960 S L18 OR L19 OR L20  
L30 0 S L29 AND L16  
L31 0 S L29 AND L15  
L32 4 S L29 AND L14  
L33 0 S L29 AND L13  
L34 0 S L29 AND L12  
L35 30 S L29 AND L11  
L36 0 S L29 AND L10  
L37 350 S L21 AND L22  
L38 36331 S L21 OR L22  
L39 4 S L38 AND L10  
L40 2 S L38 AND L11  
L41 0 S L38 AND L12  
L42 1 S L38 AND L13  
L43 46 S L38 AND L14

=> s 138 and 115

L44 1 L38 AND L15

=> d 144 1 all

L44 ANSWER 1 OF 1 CA COPYRIGHT 2001 ACS  
AN 132:288782 CA  
TI Methods and compositions for treating neurodegenerative diseases using an antagonist or inhibitor of p25  
IN Tsai, Li-Huei; Patrick, Gentry N.; Lee, Ming Sum  
PA President and Fellows of Harvard College, USA  
SO PCT Int. Appl., 54 pp.  
CODEN: PIXXD2  
DT Patent  
LA English  
IC ICM A61K038-00  
CC 1-11 (Pharmacology)  
Section cross-reference(s): 14  
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2000021550	A2	20000420	WO 1999-US24221	19991013
WO 2000021550	A3	20000727		

W: CA, JP  
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE  
PRAI US 1998-103975 19981013  
US 1999-136631 19990527  
AB The present invention relates to methods of preventing or treating neurodegenerative diseases, including Alzheimer's disease, by administering an antagonist or inhibitor of p25. In particular, the invention relates to methods of preventing or treating a neurodegenerative disease by administering a calpain antagonist or inhibitor, or a cation (e.g. Ca<sup>2+</sup>) antagonist or inhibitor, which reduces the truncation or conversion of p35 to p25. Calpeptin and ALLM, inhibitors of a calcium-activated protease (calpain), completely inhibited the conversion of p35 to p25 in calcium-treated mouse brain lysate, indicating that calpain plays an important role in the conversion process.  
ST p35 conversion p25 inhibitor neurodegenerative disease; calpain cation inhibitor nervous system agent  
IT Parkinson's disease  
(Guamanian parkinsonism-dementia; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)  
IT Nervous system  
(Huntington's chorea; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)  
IT Mental disorder  
(Pick's disease; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)  
IT Nerve  
(degeneration, corticobasal; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)  
for  
IT Mental disorder  
(dementia; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)  
IT Chromosome  
(human 17, dementia linked to; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)  
IT Anti-Alzheimer's agents

Anti-ischemic agents  
Antiparkinsonian agents  
Down's syndrome  
Nervous system agents  
Neurofibrillary tangle  
Niemann-Pick disease  
Oxidative stress, biological  
(inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Brain  
Spinal cord  
(inhibition of conversion of p35 to p25 in brain and spinal cord for treating neurodegenerative diseases)

IT Nucleic acids  
Peptides, biological studies  
RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
(inhibition of protein p25 for treating neurodegenerative diseases)

IT Brain, disease  
Heart, disease  
(ischemia; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Muscular dystrophy  
(myotonic; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Apoptosis  
(neuronal; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Dopamine agonists  
(non-ergot DE; inhibition of protein p25 for treating neurodegenerative diseases)

IT Proteins, specific or class  
RL: ADV (Adverse effect, including toxicity); ANT (Analyte); BOC (Biological occurrence); BPR (Biological process); BSU (Biological study, unclassified); MFM (Metabolic formation); ANST (Analytical study); BIOL (Biological study); FORM (Formation, nonpreparative); OCCU (Occurrence); PROC (Process)  
(p25; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Proteins, specific or class  
RL: ANT (Analyte); BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PROC (Process); USES (Uses)  
(p35; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Encephalitis  
(pan-, subacute sclerosing; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Parkinson's disease  
(postencephalic; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Phosphorylation, biological  
(protein; redn. of phosphorylation of .tau. protein by p25/cdk5 kinase for treating neurodegenerative diseases)

IT Paralysis  
(pseudobulbar; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Nervous system  
(sclerosis, lower lateral; inhibition of conversion of p35 to p25 for treating neurodegenerative diseases)

IT Antibodies

RL: BAC (Biological activity or effector, except adverse); THU  
(Therapeutic use); BIOL (Biological study); USES (Uses)  
(specific to p25 or cdk5; inhibition of protein p25 for treating  
neurodegenerative diseases)

IT Brain, disease  
(stroke; inhibition of conversion of p35 to p25 for treating  
neurodegenerative diseases)

IT Multiple sclerosis  
(therapeutic agents; inhibition of conversion of p35 to p25 for  
treating neurodegenerative diseases)

IT Prion diseases  
(with tangles; inhibition of conversion of p35 to p25 for treating  
neurodegenerative diseases)

IT Transferrins  
RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
(.tau.-transferrins, phosphorylation; redn. of phosphorylation of  
.tau.  
protein by p25/cdk5 kinase for treating neurodegenerative diseases)

IT 55123-66-5, Leupeptin 66701-25-5, E 64 79079-11-1, Calpastatin  
88191-84-8, MDL 28170 110044-82-1 110115-07-6 117591-20-5,  
Calpeptin  
158798-83-5, AK 275  
RL: BAC (Biological activity or effector, except adverse); THU  
(Therapeutic use); BIOL (Biological study); USES (Uses)  
(calpain inhibitors for treating neurodegenerative diseases)

IT 7439-95-4, Magnesium, biological studies 7440-70-2, Calcium, biological  
studies 78990-62-2, Calpain  
RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL  
(Biological study); PROC (Process)  
(calpain or cation inhibitors for treating neurodegenerative diseases)

IT 147014-96-8, Cdk5 kinase  
RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
(inhibition of deregulation of cdk5 kinase by p25 for treating  
neurodegenerative diseases)

IT 91374-21-9, Ropinirole 117630-06-5, .omega.-Conotoxin 141429-64-3, SB  
201823A 264236-80-8, MKA 01  
RL: BAC (Biological activity or effector, except adverse); THU  
(Therapeutic use); BIOL (Biological study); USES (Uses)  
(inhibition of protein p25 for treating neurodegenerative diseases)

IT 9001-66-5, Monoamine oxidase 9012-25-3, Catechol-O-methyltransferase  
RL: BSU (Biological study, unclassified); BIOL (Biological study)  
(inhibitors; inhibition of protein p25 for treating neurodegenerative  
diseases)

=> s 138 and 116

L45 0 L38 AND L16

=> s 138 and 117

L46 0 L38 AND L17

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1 11 S NEUROSERPIN  
L2 1 S PMSF  
L3 1 S APMSF  
L4 1 S L3

L5            5 S ANTIPAIN  
L6            170 S ANTITHROMBIN  
L7            11 S LEUPEPTIN  
L8            9 S DICHLOOROCOUMARIN  
L9            2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

L10            17 S L1  
               E NEUROSERPIN  
L11            728 S L2  
L12            24 S L3  
L13            343 S L5  
L14            4326 S L6  
L15            171 S L7  
L16            16 S L8  
L17            372 S L9  
               E NEUROPATHY  
L18            5490 S E3  
               E EPILEPSY  
L19            10805 S E3-E12  
               E SEIZURE  
L20            14562 S E3-E6  
               E HYPOXIA  
L21            24801 S E3  
               E STROKE  
L22            11880 S E3  
L23            0 S L17 AND L18  
L24            0 S L17 AND L19  
L25            0 S L17 AND L20  
L26            0 S L17 AND L21  
L27            0 S L17 AND L22  
L28            0 S L16 AND L18  
L29            25960 S L18 OR L19 OR L20  
L30            0 S L29 AND L16  
L31            0 S L29 AND L15  
L32            4 S L29 AND L14  
L33            0 S L29 AND L13  
L34            0 S L29 AND L12  
L35            30 S L29 AND L11  
L36            0 S L29 AND L10  
L37            350 S L21 AND L22  
L38            36331 S L21 OR L22  
L39            4 S L38 AND L10  
L40            2 S L38 AND L11  
L41            0 S L38 AND L12  
L42            1 S L38 AND L13  
L43            46 S L38 AND L14  
L44            1 S L38 AND L15  
L45            0 S L38 AND L16  
L46            0 S L38 AND L17

=> e ischemia

E1            5      ISCHELIUM/BI  
E2            112     ISCHEM/BI  
E3            9 --> ISCHEMA/BI  
E4            1      ISCHEMAI/BI  
E5            2      ISCHEMI/BI  
E6            40127    ISCHEMIA/BI  
E7            2      ISCHEMIAC/BI  
E8            1      ISCHEMIAINDUCED/BI

E9 3 ISCHEMIAL/BI  
E10 2 ISCHEMIA/BI  
E11 1 ISCHEMIAREPERFUSION/BI  
E12 44 ISCHEMIAS/BI

=> s e6

L47 40127 ISCHEMIA/BI

=> s 147 and 110

L48 3 L47 AND L10

=> d 148 1-3

L48 ANSWER 1 OF 3 CA COPYRIGHT 2001 ACS  
AN 133:187780 CA  
TI Neuroserpin reduces cerebral infarct volume and protects neurons from  
*ischemia*-induced apoptosis  
AU Yepes, Manuel; Sandkvist, Maria; Wong, Mike K. K.; Coleman, Timothy A.;  
Smith, Elizabeth; Cohan, Stanley L.; Lawrence, Daniel A.  
CS Department of Biochemistry, American Red Cross Holland Laboratory,  
Rockville, MD, 20855, USA  
SO Blood (2000), 96(2), 569-576  
CODEN: BLOOAW; ISSN: 0006-4971  
PB American Society of Hematology  
DT Journal  
LA English  
RE.CNT 70  
RE  
(1) Ahn, M; Brain Res 1999, V837, P169 CA  
(3) Benveniste, H; J Neurochem 1984, V43, P1369 CA  
(5) Calof, A; Neuron 1994, V13, P117 CA  
(7) Carroll, P; Development 1994, V120, P3173 CA  
(8) Chen, Z; Cell 1997, V91, P917 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L48 ANSWER 2 OF 3 CA COPYRIGHT 2001 ACS

AN 131:255873 CA  
TI Transforming growth factor-.beta.1 as a regulator of the serpins/t-PA  
axis

in cerebral *ischemia*  
AU Docagne, Fabian; Nicole, Olivier; Marti, Hugo H.; MacKenzie, Eric T.;  
Buisson, Alain; Vivien, Denis  
CS Universite de Caen, CNRS UMR 6551, Caen, 14074, Fr.  
SO FASEB J. (1999), 13(11), 1315-1324  
CODEN: FAJOEC; ISSN: 0892-6638

PB Federation of American Societies for Experimental Biology

DT Journal

LA English

RE.CNT 57

RE

(1) Breier, G; Development 1992, V114, P521 CA  
(2) Buisson, A; FASEB J 1998, V12, P1683 CA  
(3) Buisson, A; Neuropharmacology 1995, V34, P1081 CA  
(5) Constat, D; J Immunol 1992, V148, P1404 CA  
(6) Cunningham, D; J Cell Biochem 1989, V39, P55 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L48 ANSWER 3 OF 3 CA COPYRIGHT 2001 ACS

AN 131:139510 CA

TI Neuroserpin applications as a pharmaceutical or diagnostic agent

IN Sonderegger, Peter; Schrimpf, Sabine Petra; Kruger, Stefan Robert;  
Osterwalder, Thomas; Stockli, Esther Trudi

PA Switz.

SO PCT Int. Appl., 55 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9941381	A1	19990819	WO 1999-IB248	19990212
	W:	AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,			

TM

RW:	GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		
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AU 9921807	A1	19990830	AU 1999-21807	19990212
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PRAI US 1998-23129 19980213  
WO 1999-IB248 19990212

RE.CNT 6

RE

- (1) Coleman, T; WO 9816643 A 1998 CA
- (2) Hastings, G; THE JOURNAL OF BIOLOGICAL CHEMISTRY 1997, V272(52), P33062 CA
- (3) Incyte Pharma Inc; WO 9640922 A 1996 CA
- (4) Krueger, S; THE JOURNAL OF NEUROSCIENCE 1997, V17(23), P8984 CA
- (6) Schrimpf; GENOMICS 1997, V40(1), P55 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s 147 and 116

L49 0 L47 AND L16

=> s 147 and 115

L50 3 L47 AND L15

=> d 150 1-3

L50 ANSWER 1 OF 3 CA COPYRIGHT 2001 ACS

AN 132:288782 CA

TI Methods and compositions for treating neurodegenerative diseases using an antagonist or inhibitor of p25

IN Tsai, Li-Huei; Patrick, Gentry N.; Lee, Ming Sum

PA President and Fellows of Harvard College, USA

SO PCT Int. Appl., 54 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000021550	A2	20000420	WO 1999-US24221	19991013
	WO 2000021550	A3	20000727		
	W:	CA, JP			
	RW:	AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE			

PRAI US 1998-103975 19981013

L50 ANSWER 2 OF 3 CA COPYRIGHT 2001 ACS  
 AN 119:136686 CA  
 TI Protective effect of the protease inhibitor leupeptin against myocardial stunning  
 AU Matsumura, Yasushi; Kusuoka, Hideo; Inoue, Michitoshi; Hori, Masatsugu; Kamada, Takenobu  
 CS Med. Sch., Osaka Univ., Suita, 565, Japan  
 SO J. Cardiovasc. Pharmacol. (1993), 22(1), 135-42  
 CODEN: JCPCDT; ISSN: 0160-2446  
 DT Journal  
 LA English

L50 ANSWER 3 OF 3 CA COPYRIGHT 2001 ACS  
 AN 109:236993 CA  
 TI Carnitine-coupled pharmaceutical agents for site-specific delivery to cardiac and skeletal muscle  
 IN Stracher, Alfred; Kesner, Leo  
 PA USA  
 SO U.S., 5 pp.  
 CODEN: USXXAM  
 DT Patent  
 LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 4742081	A	19880503	US 1986-816546	19860106
	US 4866040	A	19890912	US 1987-3888	19870115
	US 5008288	A	19910416	US 1989-347361	19890504
	US 5876747	A	19990302	US 1992-912068	19920708
PRAI	US 1986-816546	19860106			
	US 1987-3888	19870115			
	US 1989-347361	19890504			
	US 1991-638948	19910109			
OS	MARPAT 109:236993				

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(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1	11 S NEUROSERPIN
L2	1 S PMSF
L3	1 S APMSF
L4	1 S L3
L5	5 S ANTIPAIN
L6	170 S ANTITHROMBIN
L7	11 S LEUPEPTIN
L8	9 S DICHLOROCOUMARIN
L9	2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

L10	17 S L1
	E NEUROSERPIN
L11	728 S L2
L12	24 S L3
L13	343 S L5
L14	4326 S L6

L15            171 S L7  
L16            16 S L8  
L17            372 S L9  
                E NEUROPATHY  
L18            5490 S E3  
                E EPILEPSY  
L19            10805 S E3-E12  
                E SEIZURE  
L20            14562 S E3-E6  
                E HYPOXIA  
L21            24801 S E3  
                E STROKE  
L22            11880 S E3  
L23            0 S L17 AND L18  
L24            0 S L17 AND L19  
L25            0 S L17 AND L20  
L26            0 S L17 AND L21  
L27            0 S L17 AND L22  
L28            0 S L16 AND L18  
L29            25960 S L18 OR L19 OR L20  
L30            0 S L29 AND L16  
L31            0 S L29 AND L15  
L32            4 S L29 AND L14  
L33            0 S L29 AND L13  
L34            0 S L29 AND L12  
L35            30 S L29 AND L11  
L36            0 S L29 AND L10  
L37            350 S L21 AND L22  
L38            36331 S L21 OR L22  
L39            4 S L38 AND L10  
L40            2 S L38 AND L11  
L41            0 S L38 AND L12  
L42            1 S L38 AND L13  
L43            46 S L38 AND L14  
L44            1 S L38 AND L15  
L45            0 S L38 AND L16  
L46            0 S L38 AND L17  
                E ISCHEMIA  
L47            40127 S E6  
L48            3 S L47 AND L10  
L49            0 S L47 AND L16  
L50            3 S L47 AND L15

=> d 147 and l11

L11 IS NOT VALID HERE  
For an explanation, enter "HELP DISPLAY".

=> s 147 and l11

L51            3 L47 AND L11

=> d 151 1-3

L51 ANSWER 1 OF 3 CA COPYRIGHT 2001 ACS  
AN 121:99561 CA  
TI Effect of phenylmethylsulfonyl fluoride (PMSF) on brain arachidonic acid  
and extracellular glutamate level in complete cerebral **ischemia**  
in rats  
AU Tanimura, Hajime  
CS Dep. Neurosurg., Kakegawa Gen. Hosp., Kakegawa, 436, Japan  
SO Brain Nerve (1994), 46(2), 153-7

CODEN: BRNED8; ISSN: 0006-8969

DT Journal  
LA Japanese

L51 ANSWER 2 OF 3 CA COPYRIGHT 2001 ACS

AN 117:168825 CA

TI A phospholipase C inhibitor ameliorates postischemic neuronal damage in rats

AU Umemura, Atsushi; Mabe, Hideo; Nagai, Hajime

CS Med. Sch., Nagoya City Univ., Nagoya, Japan

SO Stroke (Dallas) (1992), 23(8), 1163-6

CODEN: SJCCA7; ISSN: 0039-2499

DT Journal

LA English

L51 ANSWER 3 OF 3 CA COPYRIGHT 2001 ACS

AN 103:206103 CA

TI Protection by acyl-carnitines and phenylmethylsulfonyl fluoride of rat heart subjected to **ischemia** and reperfusion

AU Huelsmann, W. C.; Dubelaar, M. L.; Lamers, J. M. J.; Maccari, F.

CS Med. Fac., Erasmus Univ., Rotterdam, 3000 DR, Neth.

SO Biochim. Biophys. Acta (1985), 847(1), 62-6

CODEN: BBACAO; ISSN: 0006-3002

DT Journal

LA English

=> d 151 1 2 3 all

L51 ANSWER 1 OF 3 CA COPYRIGHT 2001 ACS

AN 121:99561 CA

TI Effect of phenylmethylsulfonyl fluoride (PMSF) on brain arachidonic acid and extracellular glutamate level in complete cerebral **ischemia** in rats

AU Tanimura, Hajime

CS Dep. Neurosurg., Kakegawa Gen. Hosp., Kakegawa, 436, Japan

SO Brain Nerve (1994), 46(2), 153-7

CODEN: BRNED8; ISSN: 0006-8969

DT Journal

LA Japanese

CC 1-11 (Pharmacology)

AB Complete cerebral **ischemia** was induced in rats by decapitation.

Tissue concns. of free arachidonic acid and extracellular levels of glutamate were measured in the striatum after the ischemic insult. PMSF inhibited arachidonic acid release during the 1st 4 min of **ischemia**. PMSF also prevented the **ischemia**-induced rise in extracellular glutamate during the 1st 4 min of **ischemia**.

Since it is known that acetylcholine inhibits glutamate release, these results suggest that PMSF inhibits acetylcholinesterase activity in the early stage of complete cerebral **ischemia** and thereby inhibits the **ischemia**-induced increase of extracellular glutamate; the inhibition of arachidonic acid release may be secondary to the inhibition of glutamate receptors rather than to an inhibition of phospholipase C activity.

ST brain **ischemia** phenylmethylsulfonyl fluoride; arachidonate metab

brain **ischemia** phenylmethylsulfonyl fluoride; glutamate metab

brain **ischemia** phenylmethylsulfonyl fluoride

IT Brain, disease

(**ischemia**, arachidonic acid and glutamic acid metab. in,  
phenylmethylsulfonyl fluoride effect on)

IT 329-98-6, Phenylmethylsulfonyl fluoride

RL: BIOL (Biological study)

(arachidonic acid and glutamic acid metab. by brain in **ischemia**  
response to)

IT 56-86-0, Glutamic acid, biological studies 506-32-1, Arachidonic acid  
RL: BPR (Biological process); BIOL (Biological study); PROC (Process)  
(metab. of, by brain in **ischemia**, phenylmethylsulfonyl  
fluoride effect on)

L51 ANSWER 2 OF 3 CA COPYRIGHT 2001 ACS

AN 117:168825 CA

TI A phospholipase C inhibitor ameliorates postischemic neuronal damage in rats

AU Umemura, Atsushi; Mabe, Hideo; Nagai, Hajime

CS Med. Sch., Nagoya City Univ., Nagoya, Japan

SO Stroke (Dallas) (1992), 23(8), 1163-6

CODEN: SJCCA7; ISSN: 0039-2499

DT Journal

LA English

CC 14-10 (Mammalian Pathological Biochemistry)

Section cross-reference(s): 1

AB Calcium-induced neuronal damage may occur in brain **ischemia**. Phospholipase C catalyzes the phosphodiester bond cleavage of phosphatidylinositol. The cleavage of phosphatidylinositol 4,5-bisphosphate by phospholipase C yields 1,4,5-inositol trisphosphate, which mediates the intracellular release of calcium, and 1,2-diacylglycerol, which is an activator of protein kinase C. The effects of phenylmethylsulfonylfluoride, a phospholipase C inhibitor, on delayed neuronal damage after a transient 20-min forebrain **ischemia** were studied in the brain hippocampal CA1 subfield in rats to assess the role of phospholipase C in postischemic neuronal damage. The neuronal d. in the CA1 subfield was detd. 7 days after reperfusion. In the vehicle treatment group, the neuronal d. was 51 cells/mm of length. The neuronal densities in the 50 and 100-mg/kg phenylmethylsulfonylfluoride pretreatment groups and the 100-mg/kg phenylmethylsulfonylfluoride posttreatment group were 99, 150, and 143 cells/mm, resp. Thus, activation of phospholipase C has an important role

in the postischemic delayed neuronal damage.

ST brain **ischemia** neuron damage phospholipase C

IT Brain, disease

(**ischemia**, phospholipase C in pathogenesis of neuron damage in)

IT 9001-86-9, Phospholipase C

RL: BIOL (Biological study)  
(in brain ischemic neuron damage pathogenesis)

IT 329-98-6, Phenylmethylsulfonylfluoride

RL: BIOL (Biological study)  
(phospholipase C inhibition by, brain neuron ischemic damage decrease by)

L51 ANSWER 3 OF 3 CA COPYRIGHT 2001 ACS

AN 103:206103 CA

TI Protection by acyl-carnitines and phenylmethylsulfonyl fluoride of rat heart subjected to **ischemia** and reperfusion

AU Huelsmann, W. C.; Dubelaar, M. L.; Lamers, J. M. J.; Maccari, F.

CS Med. Fac., Erasmus Univ., Rotterdam, 3000 DR, Neth.

SO Biochim. Biophys. Acta (1985), 847(1), 62-6

CODEN: BBACAO; ISSN: 0006-3002

DT Journal

LA English

CC 1-8 (Pharmacology)

AB Perfusion of rat hearts according to the Langendorff technique with micromolar concns. of palmitoylcarnitine [1935-18-8] or millimolar concns. of phenylmethylsulfonyl fluoride [329-98-6] protects

the heart from deterioration by reperfusion after total **ischemia**. This is based on the retention of the cytosolic enzymes detd. (lactate dehydrogenase, glycogen phosphorylase and glycogen synthase) and of myoglobin, as well as on the resumption of contractile activity. Palmitoylcarnitine, like phenylmethylsulfonyl fluoride, could protect through plasma membrane stabilization, since more hydrophilic compds. had no effect.

ST acylcarnitine phenylmethylsulfonyl fluoride heart **ischemia**

IT Hydrophilicity

(of acylcarnitines, heart protection after **ischemia** and reperfusion in relation to)

IT Heart, disease or disorder

(**ischemia**, damage from reperfusion after, acylcarnitines and phenylmethylsulfonyl fluoride protection from)

IT 329-98-6 541-14-0 541-15-1 541-15-1D, acyl derivs.

1935-18-8 17298-37-2 25243-95-2 76932-34-8

RL: BIOL (Biological study)

(heart protection from, in **ischemia** and reperfusion)

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1 11 S NEUROSERPIN  
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L5 5 S ANTIPAIN  
L6 170 S ANTITHROMBIN  
L7 11 S LEUPEPTIN  
L8 9 S DICHLOROCOUMARIN  
L9 2 S TLCK

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L10 17 S L1  
E NEUROSERPIN  
L11 728 S L2  
L12 24 S L3  
L13 343 S L5  
L14 4326 S L6  
L15 171 S L7  
L16 16 S L8  
L17 372 S L9  
E NEUROPATHY  
L18 5490 S E3  
E EPILEPSY  
L19 10805 S E3-E12  
E SEIZURE  
L20 14562 S E3-E6  
E HYPOXIA  
L21 24801 S E3  
E STROKE  
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L28            0 S L16 AND L18  
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L49            0 S L47 AND L16  
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=> s l47 and l12

L52            0 L47 AND L12

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L53            1 L47 AND L13

=> s l53 1

MISSING OPERATOR L53 1

The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> d 153 1

L53 ANSWER 1 OF 1 CA COPYRIGHT 2001 ACS  
AN 99:205847 CA  
TI Role of cellular proteinases in acute myocardial infarction. I.  
Proteolysis in nonischemic and ischemic rat myocardium and the effects of antipain, leupeptin, pepstatin and chymostatin administered in vivo  
AU Bolli, Roberto; Cannon, Richard O.; Speir, Edith; Goldstein, Robert E.; Epstein, Stephen E.  
CS Sect. Exp. Phys. Pharmacol., Natl. Heart, Lung, Blood Inst., Bethesda,  
MD,  
USA  
SO J. Am. Coll. Cardiol. (1983), 2(4), 671-80  
CODEN: JACCDI; ISSN: 0735-1097  
DT Journal  
LA English

=> d 153 1 all

L53 ANSWER 1 OF 1 CA COPYRIGHT 2001 ACS

AN 99:205847 CA  
TI Role of cellular proteinases in acute myocardial infarction. I.  
Proteolysis in nonischemic and ischemic rat myocardium and the effects of  
antipain, leupeptin, pepstatin and chymostatin administered in vivo  
AU Bolli, Roberto; Cannon, Richard O.; Speir, Edith; Goldstein, Robert E.;  
Epstein, Stephen E.  
CS Sect. Exp. Phys. Pharmacol., Natl. Heart, Lung, Blood Inst., Bethesda,  
MD,  
USA  
SO J. Am. Coll. Cardiol. (1983), 2(4), 671-80  
CODEN: JACCDI; ISSN: 0735-1097  
DT Journal  
LA English  
CC 1-8 (Pharmacology)  
Section cross-reference(s): 14  
AB To test the hypothesis that cellular proteinase [9001-92-7] contribute  
to  
ischemic myocellular death, measurements were made of tyrosine release  
(an  
index of overall proteolysis) from incubated slices of nonischemic and  
ischemic myocardium obtained at various times after coronary artery  
occlusion in rats. Proteolysis failed to increase in ischemic myocardium  
throughout the first 24 h of occlusion, when irreversible damage  
develops,  
indicating that cellular proteinases do not undergo generalized  
activation  
in this phase. The ability of the proteinase inhibitors antipain [  
37691-11-5], leupeptin, pepstatin [39324-30-6], and chymostatin  
[9076-44-2], given in vivo, to interfere with proteolysis in ischemic  
myocardium was also evaluated. Leupeptin (10 or 40 mg/kg) inhibited  
proteolysis in a dose-related fashion (-49 and -72%, resp.). Antipain  
(20  
mg/kg) decreased protein breakdown by 60%. The combination of antipain  
(20 mg/kg), leupeptin (40 mg/kg); and pepstatin 5 mg/kg) suppressed  
proteolysis almost completely at both 15 min (-88%) and at 6 h (-72%) of  
ischemia, i.e., throughout the development of irreversible injury.  
These results demonstrate that whatever proteolysis is occurring during  
acute myocardial infarction is largely mediated by cathepsins A, B, D, L  
and H and by calcium-activated neutral protease.  
ST heart infarction proteinase inhibitor  
IT Leupeptins  
RL: BIOL (Biological study)  
(proteinase inhibition by, in myocardial infarction)  
IT Proteins  
RL: BIOL (Biological study)  
(proteolysis of, in myocardial infarction, proteinase inhibition in  
relation to)  
IT Heart, disease or disorder  
(infarction, proteolysis in, proteinase inhibition in relation to)  
IT 9001-92-7  
RL: BSU (Biological study, unclassified); BIOL (Biological study)  
(inhibitors, proteolysis in myocardial infarction response to)  
IT 9076-44-2 37691-11-5 39324-30-6  
RL: BIOL (Biological study)  
(proteinase inhibition by, in myocardial infarction)  
IT 9004-08-4  
RL: BIOL (Biological study)  
(proteolysis during acute myocardial infarction in relation to)

=> d his

## FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

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 L2           1 S PMSF  
 L3           1 S APMSF  
 L4           1 S L3  
 L5           5 S ANTIPAIN  
 L6           170 S ANTITHROMBIN  
 L7           11 S LEUPEPTIN  
 L8           9 S DICHLOROCOUMARIN  
 L9           2 S TLCK

## FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

## FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

L10           17 S L1  
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 L15           171 S L7  
 L16           16 S L8  
 L17           372 S L9  
               E NEUROPATHY  
 L18           5490 S E3  
               E EPILEPSY  
 L19           10805 S E3-E12  
               E SEIZURE  
 L20           14562 S E3-E6  
               E HYPOXIA  
 L21           24801 S E3  
               E STROKE  
 L22           11880 S E3  
 L23           0 S L17 AND L18  
 L24           0 S L17 AND L19  
 L25           0 S L17 AND L20  
 L26           0 S L17 AND L21  
 L27           0 S L17 AND L22  
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 L38           36331 S L21 OR L22  
 L39           4 S L38 AND L10  
 L40           2 S L38 AND L11  
 L41           0 S L38 AND L12  
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 L47           40127 S E6  
 L48           3 S L47 AND L10  
 L49           0 S L47 AND L16

L50            3 S L47 AND L15  
L51            3 S L47 AND L11  
L52            0 S L47 AND L12  
L53            1 S L47 AND L13

=> d 147 and 114

L14 IS NOT VALID HERE  
For an explanation, enter "HELP DISPLAY".

=> s 147 and 114

L54            52 L47 AND L14

=> s his

L55            28898 HIS

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1            11 S NEUROSERPIN  
L2            1 S PMSF  
L3            1 S APMSF  
L4            1 S L3  
L5            5 S ANTIPAIN  
L6            170 S ANTITHROMBIN  
L7            11 S LEUPEPTIN  
L8            9 S DICHLOROCOUMARIN  
L9            2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

L10            17 S L1  
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L11            728 S L2  
L12            24 S L3  
L13            343 S L5  
L14            4326 S L6  
L15            171 S L7  
L16            16 S L8  
L17            372 S L9  
              E NEUROPATHY  
L18            5490 S E3  
              E EPILEPSY  
L19            10805 S E3-E12  
              E SEIZURE  
L20            14562 S E3-E6  
              E HYPOXIA  
L21            24801 S E3  
              E STROKE  
L22            11880 S E3  
              0 S L17 AND L18  
L24            0 S L17 AND L19  
L25            0 S L17 AND L20  
L26            0 S L17 AND L21  
L27            0 S L17 AND L22  
L28            0 S L16 AND L18  
L29            25960 S L18 OR L19 OR L20

L30           0 S L29 AND L16  
L31           0 S L29 AND L15  
L32           4 S L29 AND L14  
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L34           0 S L29 AND L12  
L35           30 S L29 AND L11  
L36           0 S L29 AND L10  
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L40           2 S L38 AND L11  
L41           0 S L38 AND L12  
L42           1 S L38 AND L13  
L43           46 S L38 AND L14  
L44           1 S L38 AND L15  
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L46           0 S L38 AND L17  
            E ISCHEMA  
L47           40127 S E6  
L48           3 S L47 AND L10  
L49           0 S L47 AND L16  
L50           3 S L47 AND L15  
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L54           52 S L47 AND L14  
L55           28898 S HIS

=> s 154 not 143

L56           47 L54 NOT L43

=> d 156 10-47

L56 ANSWER 10 OF 47 CA COPYRIGHT 2001 ACS  
AN 131:295340 CA  
TI Effects of a novel mac-1 inhibitor, NPC 15669, on hemostatic parameters during preconditioned myocardial infarction  
AU Serebruany, Victor L.; Yurovsky, Vladimir V.; Gurbel, Paul A.  
CS Sinai Center for Thrombosis Research, University of Maryland School of Medicine, Baltimore, MD, 21201, USA  
SO Life Sci. (1999), 65(14), 1503-1513  
CODEN: LIFSAK; ISSN: 0024-3205  
PB Elsevier Science Inc.  
DT Journal  
LA English  
RE.CNT 36  
RE  
(1) Bastida, E; Blood 1987, V70, P1437 CA  
(2) Bennett, R; Arteriosc Thromb 1993, V13, P360 CA  
(5) Cho, P; J Surg Res 1993, V54, P486 CA  
(6) Corbi, A; J Biol Chem 1988, V263, P12403 CA  
(9) D'Angelo, A; Thromb Res 1994, V75, P133 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L56 ANSWER 11 OF 47 CA COPYRIGHT 2001 ACS

AN 131:27699 CA  
TI Antithrombin III treatment improves parameters of acute inflammation in a highly histoincompatible model of rat lung allograft rejection  
AU Okada, Yoshinori; Zuo, Xiao-Jing; Marchevsky, Alberto M.; Nicolaidou, Electra; Toyoda, Mieko; Matloff, Jack M.; Jordan, Stanley C.  
CS Department of Cardiothoracic Surgery, The Cedars Sinai Medical Center

Burns and Allen Research Institute, UCLA School of Medicine, Los Angeles,  
CA, 90048, USA  
SO Transplantation (1999), 67(4), 526-528  
CODEN: TRPLAU; ISSN: 0041-1337  
PB Lippincott Williams & Wilkins  
DT Journal  
LA English  
RE.CNT 13  
RE  
(2) Altieri, D; Cell Immunol 1994, V155(2), P372 CA  
(8) Matsumura, Y; Transplantation 1995, V59, P551 CA  
(10) Okajima, K; Semin Thromb Hemost 1998, V24(1), P27 CA  
(12) Takeshita, K; Transplant Proc 1996, V28, P631 CA  
(13) Uchiba, M; Am J Physiol 1996, V270, PL921 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L56 ANSWER 12 OF 47 CA COPYRIGHT 2001 ACS  
AN 131:13277 CA  
TI Prevention of the endothelial cell injury by physiological  
anticoagulants.  
The mechanisms and therapeutic implications  
AU Okajima, Kenji  
CS Sch. Med., Kumamoto Univ., Kumamoto, 860-8556, Japan  
SO Seibutsu Shiryo Bunseki (1998), 21(4), 243-248  
CODEN: SSBUEL; ISSN: 0913-3763  
PB Seibutsu Shiryo Bunseki Kagakkai  
DT Journal; General Review  
LA Japanese

L56 ANSWER 13 OF 47 CA COPYRIGHT 2001 ACS  
AN 130:294938 CA  
TI The effect of tissue factor pathway inhibitor on hepatic ischemic  
reperfusion injury of the rat  
AU Yoshimura, Norio; Kobayashi, Yosifumi; Nakamura, Kenji; Yamagishi,  
Hisakazu; Oka, Takahiro  
CS Second Department of Surgery, Kyoto Prefectural University of Medicine,  
Kyoto City, 602, Japan  
SO Transplantation (1999), 67(1), 45-53  
CODEN: TRPLAU; ISSN: 0041-1337  
PB Lippincott Williams & Wilkins  
DT Journal  
LA English  
RE.CNT 46  
RE  
(1) Archipoff, G; Biochem J 1991, V273, P679 CA  
(2) Bajaj, M; Proc Natl Acad Sci USA 1990, V87, P8869 CA  
(4) Brand, K; Mol Cell Biol 1991, V11, P4732 CA  
(6) Brisseau, G; Blood 1995, V85, P1025 CA  
(8) Colucci, M; J Clin Invest 1983, V71, P1893 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L56 ANSWER 14 OF 47 CA COPYRIGHT 2001 ACS  
AN 130:246619 CA  
TI Treatment of severe head injury with ahyllysantifarctum  
AU Liu, Weiping; Zhang, Xiang; Yi, Shengyu; Gu, Jianwen; Song, Tao  
CS Department of Neurosurgery, 4th Military Medical University Xijing  
Hospital, Xi'an, 710033, Peop. Rep. China  
SO Disi Junyi Daxue Xuebao (1998), 19(5), 529-531  
CODEN: DJDXEG; ISSN: 1000-2790  
PB Disi Junyi Daxue Xuebao Bianjibu  
DT Journal  
LA Chinese

L56 ANSWER 15 OF 47 CA COPYRIGHT 2001 ACS  
AN 130:232253 CA  
TI Antithrombin III prevents 60 min warm intestinal **ischemia**  
reperfusion injury in rats  
AU Ozden, Akin; Tetik, Cihat; Bilgihan, Ayse; Calli, Nese; Bostanci, Birol;  
Yis, Ozgur; Duzcan, Ender  
CS Medical School, Dep. Surgery, Pamukkale Univ., Denizli, Turk.  
SO Res. Exp. Med. (1999), 198(5), 237-246  
CODEN: REXMAS; ISSN: 0300-9130  
PB Springer-Verlag  
DT Journal  
LA English  
RE.CNT 30  
RE  
(7) Grisham, M; Am J Physiol 1986, V251, PG567 CA  
(9) Hisama, N; Dig Dis Sci 1996, V41, P1481 CA  
(10) Kirchhofer, D; Blood 1993, V81, P2050 CA  
(11) Li, X; Transplantation Proc 1994, V26, P2423 CA  
(12) Matsutani, T; J Surg Res 1998, V79, P158 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L56 ANSWER 16 OF 47 CA COPYRIGHT 2001 ACS  
AN 130:176992 CA  
TI The anti-inflammatory properties of antithrombin III: new therapeutic  
implications  
AU Okajima, Kenji; Uchiba, Mitsuhiro  
CS Department of Laboratory Medicine, Kumamoto University School of  
Medicine,  
Kumamoto, 860, Japan  
SO Semin. Thromb. Hemostasis (1998), 24(1), 27-32  
CODEN: STHMBV; ISSN: 0094-6176  
PB Thieme Medical Publishers, Inc.  
DT Journal; General Review  
LA English  
RE.CNT 35  
RE  
(1) Atalla, S; Transplantation 1985, V40, P584 CA  
(4) Coalson, J; Circ Shock 1978, V5, P423 CA  
(5) Eisenhut, T; Immunopharmacol 1993, V26, P259 CA  
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(9) Harada, N; The Immune Consequence of Trauma, Shock and Sepsis: Mechanisms  
and Therapeutic Approaches 1997, P625 CA  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L56 ANSWER 17 OF 47 CA COPYRIGHT 2001 ACS  
AN 130:61081 CA  
TI Compositions for treating and preventing arterial thrombosis and use of a  
factor Xa inhibitor alone or combined with a platelet aggregation  
inhibitor  
IN Bernat, Andre; Herbert, Jean-Marc; Petitou, Maurice; Van Amsterdam,  
Ronald  
PA Sanofi, Fr.; Akzo Nobel N.V.  
SO PCT Int. Appl., 90 pp.  
CODEN: PIXXD2  
DT Patent  
LA French  
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 9856365	A1	19981217	WO 1998-FR1172	19980609
W:	AU, BR, BY, CA, CN, CZ, EE, HU, ID, IL, IS, JP, KR, LK, LT, LV, MX, NO, NZ, PL, RU, SG, SI, SK, TR, UA, US, VN, YU			
RW:	AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,			

PT, SE				
FR 2764511	A1	19981218	FR 1997-7368	19970613
FR 2764511	B1	20000908	AU 1998-79246	19980609
AU 9879246	A1	19981230		
AU 728826	B2	20010118		
EP 986376	A1	20000322	EP 1998-929521	19980609
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
BR 9810520	A	20000919	BR 1998-10520	19980609
ZA 9805137	A	19990107	ZA 1998-5137	19980612
NO 9906137	A	20000214	NO 1999-6137	19991210
PRAI FR 1997-7368		19970613		
WO 1998-FR1172		19980609		

RE.CNT 21

RE

- (1) Bernat; Fibrinolysis 1996, V10(3), P151 CA
  - (2) Cadroy, Y; Thrombosis and Haemostasis V70(4), P631 CA
  - (3) Choay; EP 0138632 A 1985 CA
  - (4) Daiichi Seiyaku Co; EP 0540051 A 1993 CA
  - (7) Herault; Blood Coagul Fibrinolysis 1997, V8(3), P206 CA
- ALL CITATIONS AVAILABLE IN THE RE FORMAT

L56 ANSWER 18 OF 47 CA COPYRIGHT 2001 ACS

AN 129:342251 CA

TI Lipoprotein(a) level does not predict restenosis after percutaneous transluminal coronary angioplasty

AU Alaiyah, Poonam; Hoffman, Carol J.; Korlipara, Giridhar; Neuroth, Arlene; Dervan, John P.; Lawson, William E.; Hultin, Mae B.

CS Department of Medicine, State University at New York at Stony Brook, NY, USA

SO Arterioscler., Thromb., Vasc. Biol. (1998), 18(8), 1281-1286

CODEN: ATVBFA; ISSN: 1079-5642

PB Lippincott Williams & Wilkins

DT Journal

LA English

L56 ANSWER 19 OF 47 CA COPYRIGHT 2001 ACS

AN 129:270382 CA

TI Effect of low-dose heparin on fibrinogen levels in patients with chronic ischemic heart disease

AU Prisco, D.; Paniccia, R.; Bandinelli, B.; Gori, A. M.; Attanasio, M.; Giusti, B.; Comeglio, M.; Abbate, R.; Gensini, G. F.; Neri Serneri, G. G.

CS Inst. Clinica Medica Generale Cardiologia, Univ. Florence, Florence, I-50134, Italy

SO Int. J. Clin. Lab. Res. (1998), 28(3), 170-173

CODEN: ICLREA; ISSN: 0940-5437

PB Springer-Verlag

DT Journal

LA English

L56 ANSWER 20 OF 47 CA COPYRIGHT 2001 ACS

AN 129:270298 CA

TI Antithrombin III attenuates **ischemia/reperfusion** injury of rat liver by inhibiting leukocyte activation

AU Harada, N.; Okajima, K.; Kushimoto, S.; Isobe, H.; Uchiba, M.; Murakami, K.; Tanaka, K.; Okabe, H.

CS Department of Emergency and Critical Care Medicine, Fukuoka University, Fukuoka, Japan

SO Immune Consequences Trauma, Shock Sepsis, Int. Congr., 4th (1997), 625-630. Editor(s): Faist, Eugen. Publisher: Monduzzi Editore, Bologna, Italy.

CODEN: 66MUAY

DT Conference

LA English

L56 ANSWER 21 OF 47 CA COPYRIGHT 2001 ACS  
AN 129:183693 CA  
TI Antithrombin and **ischemia/reperfusion**  
AU Woodman, Richard C.; Ostrovsky, Lena; Teoh, Diane; Payne, Derrice; Poon, Man-Chiu; Kubes, Paul  
CS Immunology Research Group, University of Calgary, Calgary, AB, T2N 4N1, Can.  
SO Blood Coagulation Fibrinolysis (1998), 9(Suppl. 2, Potential Applications of Antithrombin Concentrate in Systemic Inflammatory Disorders), S7-S15  
CODEN: BLFIE7; ISSN: 0957-5235  
PB Lippincott-Raven Publishers  
DT Journal; General Review  
LA English

L56 ANSWER 22 OF 47 CA COPYRIGHT 2001 ACS  
AN 129:131043 CA  
TI New aspects of the antiinflammatory effect of AT III. Reduction of the reperfusion damage after warm hepatic **ischemia**  
AU Maksan, Sasa-Marcel; Gebhard, M. M.; Maksan, M.-O.; Herfarth, C.; Klar, E.  
CS Chirurgische Klinik, Abteilung Experimentelle Chirurgie, Universitaet Heidelberg, Heidelberg, D-69120, Germany  
SO Chir. Forum Exp. Klin. Forsch. (1998) 383-385  
CODEN: CFEKA7; ISSN: 0303-6227  
PB Springer-Verlag  
DT Journal  
LA German

L56 ANSWER 23 OF 47 CA COPYRIGHT 2001 ACS  
AN 128:191142 CA  
TI Elevated tissue factor and tissue factor pathway inhibitor circulating levels in ischemic heart disease patients  
AU Falciani, Michela; Gori, Anna Maria; Fedi, Sandra; Chiarugi, Ludia; Simonetti, Ignazio; Dabizzi, Roberto Piero; Prisco, Domenico; Pepe, Guglielmina; Abbate, Rosanna; Gensini, Gian Franco; Neri Serneri, Gian Gastone  
CS Istituto Clinica Medica Generale Cardiologia, University Florence, Florence, I-50134, Italy  
SO Thromb. Haemostasis (1998), 79(3), 495-499  
CODEN: THHADQ; ISSN: 0340-6245  
PB F. K. Schattauer Verlagsgesellschaft mbH  
DT Journal  
LA English

L56 ANSWER 24 OF 47 CA COPYRIGHT 2001 ACS  
AN 127:326191 CA  
TI Antithrombin III prevents and rapidly reverses leukocyte recruitment in **ischemia/reperfusion**  
AU Ostrovsky, Lena; Woodman, Richard C.; Payne, Derrice; Teoh, Diane; Kubes, Paul  
CS Department of Physiology and Biophysics, University of Calgary, Calgary, AB, T2N 4N1, Can.  
SO Circulation (1997), 96(7), 2302-2310  
CODEN: CIRCAZ; ISSN: 0009-7322  
PB American Heart Association  
DT Journal  
LA English

L56 ANSWER 25 OF 47 CA COPYRIGHT 2001 ACS  
AN 127:189106 CA  
TI Sinusoidal flow block after warm **ischemia** in rats with

AU diet-induced fatty liver  
Hakamada, Kenichi; Sasaki, Mutsuo; Takahashi, Katsuro; Umehara, Yutaka;  
Konn, Mitsuru  
CS Second Department of Surgery, Hirosaki University School of Medicine,  
Hirosaki, 036, Japan  
SO J. Surg. Res. (1997), 70(1), 12-20  
CODEN: JSGRA2; ISSN: 0022-4804  
PB Academic  
DT Journal  
LA English

L56 ANSWER 26 OF 47 CA COPYRIGHT 2001 ACS

AN 127:39825 CA

TI Human antithrombin III for treatment of ischemic reperfusion-related  
liver

damage and compositions containing human antithrombin III

IN Okajima, Kenji; Kushimoto, Shigeki

PA Green Cross Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI JP 09110718	A2	19970428	JP 1995-266829	19951016

L56 ANSWER 27 OF 47 CA COPYRIGHT 2001 ACS

AN 126:809 CA

TI Serial changes of natural antithrombotics during myocardial  
ischemia-reperfusion in swine. Effects of magnesium, diltiazem,  
and a novel Mac-1 inhibitor

AU Serebruany, V. L.; Herzog, W. R.; Gurbel, P. A.

CS Union Memorial Hospital, Heart Associates Research and Education  
Foundation, Baltimore, MD, 21218, USA

SO Blood Coagulation Fibrinolysis (1996), 7(6), 632-640

CODEN: BLFIE7; ISSN: 0957-5235

PB Rapid Science Publishers

DT Journal

LA English

L56 ANSWER 28 OF 47 CA COPYRIGHT 2001 ACS

AN 125:339091 CA

TI Pharmaceutical compositions containing human antithrombin-III for  
shock-induced gastric mucosa disorders

IN Okajima, Kenji; Kushimoto, Shigeki

PA Green Cross Corp., Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	-----	-----	-----	-----
PI JP 08245419	A2	19960924	JP 1995-45748	19950306

L56 ANSWER 29 OF 47 CA COPYRIGHT 2001 ACS

AN 125:292706 CA

TI Mac-1 inhibitor affects certain hemostatic parameters during myocardial  
stunning in swine

AU Serebruany, Victor L.; Solomon, Scott R.; Edenbaum, Lisa R.; Herzog,  
William R.; Gurbel, Paul A.

CS Heart Associated Res. Education Foundation, Union Memorial Hospital,

SO Baltimore, MD, 21218, USA  
Pharmacology (1996), 53(2), 87-97  
CODEN: PHMGBN; ISSN: 0031-7012  
DT Journal  
LA English

L56 ANSWER 30 OF 47 CA COPYRIGHT 2001 ACS  
AN 125:265930 CA  
TI The effects of breviscapin on AT-III activity, tPA and PAI in dogs during acute myocardial ischemia  
AU Sheng, Jing; Xu, Jimin; Yang, Juxian; Huang, Zhenhua; Wang, Jian; Xu, Weiren  
CS Ninth People's Hospital, SSMU, Shanghai, 200011, Peop. Rep. China  
SO J. Shanghai Second Med. Univ. (1995), 9(2), 69-73  
CODEN: JSSUE7; ISSN: 1001-6686  
DT Journal  
LA English

L56 ANSWER 31 OF 47 CA COPYRIGHT 2001 ACS  
AN 125:158640 CA  
TI New clinical uses for human-derived antithrombin III  
IN Okajima, Kenji; Taoka, Juji  
PA Green Cross Corp, Japan  
SO Jpn. Kokai Tokkyo Koho, 5 pp.  
CODEN: JKXXAF  
DT Patent  
LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 08169845	A2	19960702	JP 1995-7731	19950120
PRAI JP 1994-5131		19940121		
JP 1994-50813		19940322		
JP 1994-256508		19941021		

L56 ANSWER 32 OF 47 CA COPYRIGHT 2001 ACS  
AN 125:104730 CA  
TI The effects of Breviscapine on AT-III activity, tPA and PAI in dogs during acute myocardial ischemia  
AU Zhao, Peiqi; Xu, Jimin; Sheng, Jing; Yang, Juxiang; Huang, Zhenghua; Wang, Jian  
CS Cardiovascular Research Division, Ninth People's Hospital, Shanghai Second Medical University, Shanghai, 200011, Peop. Rep. China  
SO Shanghai Dier Yike Daxue Xuebao (1996), 16(1), 26-28  
CODEN: SDDXE3; ISSN: 0258-5898  
DT Journal  
LA Chinese

L56 ANSWER 33 OF 47 CA COPYRIGHT 2001 ACS  
AN 124:114295 CA  
TI Abnormalities in oxygenation, coagulation, and fibrinolysis in colonic blood of horses with experimentally induced strangulation obstruction  
AU Kawcak, C. E.; Baxter, G. M.; Getzy, D. M.; Stashak, T. S.; Chapman, P. L.  
CS College of Natural Sciences, Colorado State University, Fort Collins, CO, 80523, USA  
SO Am. J. Vet. Res. (1995), Volume Date 1995, 56(12), 1642-50  
CODEN: AJVRAH; ISSN: 0002-9645  
DT Journal  
LA English

L56 ANSWER 34 OF 47 CA COPYRIGHT 2001 ACS  
AN 121:252796 CA  
TI Prothrombin fragment F1 + 2: correlations with cardiovascular risk factors  
AU Rugman, F. P.; Jenkins, J. A.; Duguid, J. K.; Maggs, P. Bolton; Hay, C. R.

M.  
CS University Department of Haematology, Royal Liverpool Hospital, Liverpool,  
L7 3BX, UK  
SO Blood Coagulation Fibrinolysis (1994), 5(3), 335-40  
CODEN: BLFIE7; ISSN: 0957-5235  
DT Journal  
LA English

L56 ANSWER 35 OF 47 CA COPYRIGHT 2001 ACS  
AN 121:103160 CA  
TI Artificial ultraviolet whole-body radiation does not modify serum lipoprotein, plasma fibrinogen, plasminogen or antithrombin III concentrations in post-myocardial infarction patients  
AU Clark, Peter; Cockburn, Forrester; Cowan, Robert A.; Czapla, Krystyna; Dunnigan, Matthew G.; Farish, Elizabeth; Hughes, Elaine  
CS Medical Division, Departments of Biochemistry and Physiotherapy, Stobhill General Hospital, Glasgow, G21 3UW, UK  
SO Atherosclerosis (Shannon, Irel.) (1994), 107(1), 65-9  
CODEN: ATHSBL; ISSN: 0021-9150  
DT Journal  
LA English

L56 ANSWER 36 OF 47 CA COPYRIGHT 2001 ACS  
AN 121:80020 CA  
TI Protein S and protein C anticoagulant activity in acute and chronic cardiac ischemic syndromes. Relationship to inflammation, complement activation and in vivo thrombin activity  
AU D'Angelo, Armando; Gerosa, Stefano; Digano, Silvana; Angelo, Silvana Vigano; Mailhac, Alessandra; Colombo, Alessandro; Agazzi, Alberto; Mazzola, Giuseppina; Chierchia, Sergio  
CS Coagulation Service and Department of Cardiology, I.R.C.C.S. H S.Raffaele,  
Milan, 20132, Italy  
SO Thromb. Res. (1994), 75(2), 133-42  
CODEN: THBRAA; ISSN: 0049-3848  
DT Journal  
LA English

L56 ANSWER 37 OF 47 CA COPYRIGHT 2001 ACS  
AN 118:165910 CA  
TI Evaluation of endothelial anticoagulant function with venoocclusive test  
AU Zateyshchikov, D. A.; Dobrovolsky, A. B.; Averkov, O. V.; Storozhilova, A.  
N.; Panchenko, E. P.; Bonnet, J.; Grattsiansky, N. A.  
CS Cent. Atherosclerosis, Inst. Phys. Chem. Med., Moscow, Russia  
SO Byull. Eksp. Biol. Med. (1992), 114(12), 605-8  
CODEN: BEBMAE; ISSN: 0365-9615  
DT Journal  
LA Russian

L56 ANSWER 38 OF 47 CA COPYRIGHT 2001 ACS  
AN 112:116350 CA  
TI Antithrombin III and procoagulant activity: sex differences and effects of the menopause  
AU Meade, T. W.; Dyer, Sandra; Howarth, D. J.; Imeson, J. D.; Stirling,

Yvonne  
CS MRC Epidemiol. Med. Care Unit, Northwick Park Hosp., Harrow/Middlesex, UK  
SO Br. J. Haematol. (1990), 74(1), 77-81  
CODEN: BJHEAL; ISSN: 0007-1048

DT Journal  
LA English

L56 ANSWER 39 OF 47 CA COPYRIGHT 2001 ACS  
AN 110:229535 CA

TI Serum lipids and platelet functions in ischemic cerebrovascular diseases  
AU Takeuchi, Megumi; Uchiyama, Sinichiro; Kobayashi, Itsuro; Takemiya, Toshiko; Maruyama, Shoichi  
CS Neurol. Inst., Tokyo Women's Med. Coll., Tokyo, 162, Japan  
SO Tokyo Joshi Ika Daigaku Zasshi (1989), 59(3), 177-83  
CODEN: TJIZAF; ISSN: 0040-9022  
DT Journal  
LA Japanese

L56 ANSWER 40 OF 47 CA COPYRIGHT 2001 ACS  
AN 110:205380 CA

TI Effect of intracisternal antithrombin III on subarachnoid hemorrhage-induced arterial narrowing  
AU Vollmer, Dennis G.; Hongo, Kazuhiro; Kassell, Neal F.; Ogawa, Hisayuki; Tsukahara, Tetsuya; Lehman, R. Michael  
CS Sch. Med., Univ. Virginia, Charlottesville, VA, USA  
SO Hum. Pathol. (1989), 20(4), 599-604  
CODEN: HPCQA4; ISSN: 0046-8177  
DT Journal  
LA English

L56 ANSWER 41 OF 47 CA COPYRIGHT 2001 ACS  
AN 109:1009 CA

TI Relationship between sex hormones and hemostatic factors in healthy middle-aged men  
AU Bonithon-Kopp, Claire; Scarabin, Pierre Yves; Bara, Lucienne; Castanier, Michel; Jacqueson, Alain; Roger, Marc  
CS Hop. Broussais, Paris, 75674, Fr.  
SO Atherosclerosis (Shannon, Ireln.) (1988), 71(1), 71-6  
CODEN: ATHSBL; ISSN: 0021-9150  
DT Journal  
LA English

L56 ANSWER 42 OF 47 CA COPYRIGHT 2001 ACS  
AN 95:4832 CA

TI Studies on the clinical significance of antithrombin III with special reference to its metabolism  
AU Okuda, Seisuke  
CS Second Dep. Intern. Med., Kyoto Prefect. Univ. Med., Kyoto, Japan  
SO Kyoto-furitsu Ika Daigaku Zasshi (1981), 90(3), 247-64  
CODEN: KFIZAO  
DT Journal  
LA Japanese

L56 ANSWER 43 OF 47 CA COPYRIGHT 2001 ACS  
AN 93:218538 CA

TI Biosynthesis of antithrombin III (AT III) in rat  
AU Okuda, Seisuke; Okajima, Yasushi; Kawamura, Tsunehiro; Urano, Sumio; Nishizawa, Akihiko; Kitani, Teruo; Watada, Mitsuro; Nakagawa, Masao; Ijichi, Hamao  
CS 2nd Dep. Med., Kyoto Prefect. Univ. Med., Kyoto, Japan  
SO Ketsueki to Myakkan (1980), 11(1), 121-4  
CODEN: KTMYA3; ISSN: 0386-9717  
DT Journal

LA Japanese

L56 ANSWER 44 OF 47 CA COPYRIGHT 2001 ACS

AN 93:203085 CA

TI Hemostatic variables in vegetarians and non-vegetarians

AU Haines, A. P.; Chakrabarti, R.; Fisher, Diana; Meade, T. W.; North, W. R.  
S.; Stirling, Yvonne

CS MRC Epidemiol. Med. Care Unit, Northwick Park Hosp., Harrow/Middlesex,

HA1 3UJ, Engl.

SO Thromb. Res. (1980), 19(1-2), 139-48

CODEN: THBRAA; ISSN: 0049-3848

DT Journal

LA English

L56 ANSWER 45 OF 47 CA COPYRIGHT 2001 ACS

AN 92:196003 CA

TI Role of antithrombin III in experimental and clinical states of increased  
thrombin generation in the blood

AU Pastorova, V. E.

CS USSR

SO Vestn. Mosk. Univ., Ser. 16: Biol. (1980), (1), 18-24

CODEN: VMUBDF

DT Journal

LA Russian

L56 ANSWER 46 OF 47 CA COPYRIGHT 2001 ACS

AN 91:104211 CA

TI The use of chromogenic substrates for the determination of kallikrein and  
other serine proteases in plasma and synovial fluid in man

AU Lewis, David H.; Bengtsson, Maj Britt; Liljedahl, Sten Otto; Larsson,  
Joergen

CS Clin. Res. Cent., Univ. Hosp., Linkoeping, S-581 85, Swed.

SO Adv. Biosci. (1979), 17(Curr. Concepts Kinin Res.), 163-71

CODEN: AVBIB9; ISSN: 0065-3446

DT Journal

LA English

L56 ANSWER 47 OF 47 CA COPYRIGHT 2001 ACS

AN 87:36920 CA

TI Content of antithrombin III, fibrinogen and its degradation products,  
soluble fibrin in the blood and components of the fibrinolytic system in  
the urine in patients with ischemic heart disease

AU Panchenko, V. M.; Andreenko, G. V.; Podorol'skaya, A. V.; Bazgadze, V. M.

CS Mosk. Gos. Univ., Moscow, USSR

SO Klin. Med. (Moscow) (1977), 55(1), 25-31

CODEN: KLMIAZ

DT Journal

LA Russian

=> d 156 31 24 16 14 all

L56 ANSWER 31 OF 47 CA COPYRIGHT 2001 ACS

AN 125:158640 CA

TI New clinical uses for human-derived antithrombin III

IN Okajima, Kenji; Taoka, Juji

PA Green Cross Corp, Japan

SO Jpn. Kokai Tokkyo Koho, 5 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

IC ICM A61K038-55  
ICS A61K038-55  
CC 1-11 (Pharmacology)  
Section cross-reference(s): 63

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 08169845	A2	19960702	JP 1995-7731	19950120
PRAI	JP 1994-5131		19940121		
	JP 1994-50813		19940322		
	JP 1994-256508		19941021		
AB	Human-derived antithrombin III is claimed for prevention and treatment of motor functional disturbance, tissue injury, spinal injury, and spinal <b>ischemia</b> . The antithrombin III can be formulated into any dosage forms. Thus, i.v. injections contg. human-derived antithrombin III were prepd., and their efficacy were tested in rat models.				
ST	antithrombin III spinal injury <b>ischemia</b> ; New antithrombin III				
IT	Animal tissue				
	Spinal cord (disease, injury, new clin. uses for human-derived antithrombin III)				
IT	Spinal cord (disease, <b>ischemia</b> , new clin. uses for human-derived antithrombin III)				
IT	Pharmaceutical dosage forms (injections, i.v., new clin. uses for human-derived antithrombin III)				
IT	Nerve, disease (motor, new clin. uses for human-derived antithrombin III)				
IT	9000-94-6, Antithrombin III				
	RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (new clin. uses for human-derived antithrombin III)				

L56 ANSWER 24 OF 47 CA COPYRIGHT 2001 ACS

AN 127:326191 CA

TI Antithrombin III prevents and rapidly reverses leukocyte recruitment in **ischemia/reperfusion**

AU Ostrovsky, Lena; Woodman, Richard C.; Payne, Derrice; Teoh, Diane; Kubes, Paul

CS Department of Physiology and Biophysics, University of Calgary, Calgary, AB, T2N 4N1, Can.

SO Circulation (1997), 96(7), 2302-2310

CODEN: CIRCAZ; ISSN: 0009-7322

PB American Heart Association

DT Journal

LA English

CC 1-7 (Pharmacology)

Section cross-reference(s): 15

AB P-selectin has recently been shown to be essential for leukocyte rolling after the reperfusion of ischemic mesentery. However, the mediators responsible for neutrophil rolling in ischemic microvessels remain entirely unclear. Intravital microscopy was used to examine leukocyte kinetics in a feline mesentery **ischemia/reperfusion** model. Sixty minutes of **ischemia** followed by reperfusion caused a profound increase in leukocyte rolling and adhesion. Pretreatment with the endogenous antithrombotic agent antithrombin III (ATIII) infused as a bolus (250 U/kg) reduced neutrophil rolling and adhesion to preischemic levels during reperfusion. No effect was seen with heat-inactive ATIII. Importantly, ATIII posttreatment also significantly reduced neutrophil rolling and adhesion during reperfusion, suggesting that ATIII can reverse the leukocyte recruitment response induced by **ischemia/reperfusion**. Vascular permeability was also reduced by 50% after ATIII administration. To det. whether ATIII could reverse thrombin-induced rolling directly, neutrophil rolling was performed on human endothelium in

flow chambers. Indeed, thrombin-induced rolling, but not histamine-induced rolling, could be rapidly reversed with ATIII on endothelium, suggesting that ATIII affects thrombin rather than directly affecting neutrophils or the endothelium. This study demonstrates for

the

first time that thrombin plays an important role in **ischemia**-induced leukocyte rolling and adhesion and that ATIII can be used therapeutically postreperfusion to attenuate the leukocyte recruitment response in inflammation without the nonspecific effects assocd. with anti-adhesion mol. therapy.

ST antithrombin III leukocyte recruitment **ischemia** reperfusion

IT Blood flow

Inflammation

**Ischemia**

Leukocyte

Leukocyte rolling

Microvessel

Neutrophil

Neutrophil adhesion

Reperfusion injury

Vascular endothelium

Vascular permeability

(antithrombin III prevents and rapidly reverses leukocyte recruitment in **ischemia/reperfusion**)

IT Peritoneal diseases

(mesenteric **ischemia**; antithrombin III prevents and rapidly reverses leukocyte recruitment in **ischemia/reperfusion**)

IT **Ischemia**

(mesenteric; antithrombin III prevents and rapidly reverses leukocyte recruitment in **ischemia/reperfusion**)

IT 9002-04-4, Thrombin

RL: ADV (Adverse effect, including toxicity); BAC (Biological activity or effector, except adverse); BIOL (Biological study)

(antithrombin III prevents and rapidly reverses leukocyte recruitment in **ischemia/reperfusion**)

IT 9000-94-6, Antithrombin

RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(antithrombin III prevents and rapidly reverses leukocyte recruitment in **ischemia/reperfusion**)

L56 ANSWER 16 OF 47 CA COPYRIGHT 2001 ACS

AN 130:176992 CA

TI The anti-inflammatory properties of antithrombin III: new therapeutic implications

AU Okajima, Kenji; Uchiba, Mitsuhiro

CS Department of Laboratory Medicine, Kumamoto University School of Medicine,

Kumamoto, 860, Japan

SO Semin. Thromb. Hemostasis (1998), 24(1), 27-32

CODEN: STHMBV; ISSN: 0094-6176

PB Thieme Medical Publishers, Inc.

DT Journal; General Review

LA English

CC 1-0 (Pharmacology)

AB A review with 35 refs. Antithrombin III (AT III) supplementation has proven to be effective in the treatment of disseminated intravascular coagulation. Administration of AT III is also useful for prevention of organ failure in animals challenged with endotoxin or bacteria and it increases the survival rate of such animals. Since inhibition of coagulation abnormalities failed to prevent organ failure in animals given

bacteria, AT III may exert a therapeutic effect independent of its

anticoagulant effect. This therapeutic mechanism of AT III has been explored using an animal model of septicemia. AT III prevented pulmonary vascular injury by inhibiting leukocyte activation in rats given endotoxin. This effect is mediated by the promotion of endothelial release of prostacyclin which inhibits leukocyte activation. Interaction of AT III with heparin-like glycosaminoglycans (GAGs) on the endothelial cell surface appears to be important for this effect. Heparin inhibits these therapeutic effects of AT III by preventing AT III from interacting with the cell surface heparin-like GAGs. This activity of AT III may explain why AT III prevents organ failure as well as coagulation abnormalities in patients with sepsis. This antiinflammatory activity of AT III may be useful for the treatment of organ failure such as in **ischemia/reperfusion**-induced organ dysfunction, in which activated leukocytes play a crit. role.

ST review antiinflammatory antithrombin therapeutic  
IT Anti-inflammatory drugs  
    (anti-inflammatory activity of antithrombin and therapeutic implications)  
IT **9000-94-6, Antithrombin**  
RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
    (anti-inflammatory activity of antithrombin and therapeutic implications)

RE.CNT 35

RE

- (1) Atalla, S; Transplantation 1985, V40, P584 CA
- (2) Buller, H; Am J Med 1989, V87(Suppl 3B), P44S
- (3) Clotta, F; Am J Pathol 1994, V144, P975
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- (5) Eisenhut, T; Immunopharmacol 1993, V26, P259 CA
- (6) Emerson, T; Am J Med 1989, V87(Suppl 3B), P27S
- (7) Fourrier, F; Chest 1993, V194, P882
- (8) Gordon, J; Br J Pharmacol 1983, V80, P179 CA
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- (11) Jaeschke, H; FASEB J 1990, V4, P3355 MEDLINE
- (12) Jochum, M; Semin Hematol 1995, V32(Suppl 2), P19
- (13) Kainoh, M; Biochem Pharmacol 1990, V39, P477 CA
- (14) Kim, Y; Transplantation 1994, V58, P875 CA
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- (21) Spannagel, M; Thromb Res 1981, V61, P1
- (22) St John, R; Chest 1993, V103, P932 MEDLINE
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- (25) Tateson, J; Prostaglandins 1977, V13, P389 CA
- (26) Taylor, F; Blood 1991, V78, P364
- (27) Taylor, F; Circ Shock 1988, V26, P227 CA
- (28) Triantaphyllopoulos, D; Thromb Haemostas 1984, V51, P232 CA
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- (30) Uchiba, M; Thromb Res 1995, V80, P201 CA
- (31) Uchiba, M; to be published in Thromb Res 1998
- (32) Vinazzer, H; Semin Thromb Hemostas 1989, V15, P347 MEDLINE
- (33) Walker, R; Adv Shock Res 1982, V7, P125 CA
- (34) Weiss, S; Blood 1980, V55, P1020 CA
- (35) Yamauchi, T; Biochem Biophys Res Commun 1989, V163, P1404 CA

AN 130:246619 CA  
TI Treatment of severe head injury with ahylysantifarctum  
AU Liu, Weiping; Zhang, Xiang; Yi, Shengyu; Gu, Jianwen; Song, Tao  
CS Department of Neurosurgery, 4th Military Medical University Xijing  
Hospital, Xi'an, 710033, Peop. Rep. China  
SO Disi Junyi Daxue Xuebao (1998), 19(5), 529-531  
CODEN: DJDXEG; ISSN: 1000-2790  
PB Disi Junyi Daxue Xuebao Bianjibu  
DT Journal  
LA Chinese  
CC 1-8 (Pharmacology)  
AB Ahylysantifarctum was an antithrombin enzyme extd. from the Pallas pit viper snake venom. Sixty-nine patients with severe head trauma were randomly allocated to receive ahylysantifarctum therapy, traditional therapy, and nimodipine therapy sep. to search a better treatment of early cerebral microcirculation disturbances. There were significantly increased blood viscosity, hematocrit, and fibrinogen levels after severe head trauma. The patients' CSF exhibited metabolic acidosis, pH was 7.31.+-0.07, PCO<sub>2</sub> was 6.36.+-1.13 kPa, HCO<sub>3</sub><sup>-</sup> was 21.97.+-10 mmol/L; and the decrease of pH was related with the severity of the condition. There were significant decrease in the blood viscosity, hematocrit, fibrinogen levels in the ahylysantifarctum therapy group, P<0.05; and the CSF environment was improved, the pH raised to 7.39.+-0.09, P< 0.05;  
PCO<sub>2</sub> reduced to 5.59.+-1.38 kPa, HCO<sub>3</sub><sup>-</sup> increased to 24.09.+-1.92 mmol/L, P< 0.01; and the intracranial pressure reduced and cerebral perfusion pressure raised in some extent, P< 0.01. The results suggest the Pallas pit viper snake venom antithrombin enzyme ahylysantifarctum therapy solves the problem of impairment of cerebral microcirculation and ischemia after severe trauma, thus inducing beneficial effects in treatment of head injury.  
ST ahylysantifarctum antithrombotic head trauma; antithrombin head trauma ischemia  
IT Snake venoms  
(Pallas pit viper; treatment of severe head injury with antithrombin enzyme ahylysantifarctum)  
IT Cerebral blood flow  
(micro-; treatment of severe head injury with antithrombin enzyme ahylysantifarctum)  
IT Head injury  
(trauma; treatment of severe head injury with antithrombin enzyme ahylysantifarctum)  
IT Anti-ischemic agents  
Antithrombotics  
Cerebral ischemia  
(treatment of severe head injury with antithrombin enzyme ahylysantifarctum)  
IT 9000-94-6, Antithrombin  
RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
(treatment of severe head injury with antithrombin enzyme ahylysantifarctum)

=> d his

(FILE 'HOME' ENTERED AT 15:24:51 ON 12 APR 2001)

FILE 'REGISTRY' ENTERED AT 15:24:57 ON 12 APR 2001

L1 11 S NEUROSERPIN

L2 1 S PMSF  
L3 1 S APMSF  
L4 1 S L3  
L5 5 S ANTIPAIN  
L6 170 S ANTITHROMBIN  
L7 11 S LEUPEPTIN  
L8 9 S DICHLOROCOUMARIN  
L9 2 S TLCK

FILE 'REGISTRY' ENTERED AT 15:30:01 ON 12 APR 2001

FILE 'CA' ENTERED AT 15:30:55 ON 12 APR 2001

L10 17 S L1  
E NEUROSERPIN  
L11 728 S L2  
L12 24 S L3  
L13 343 S L5  
L14 4326 S L6  
L15 171 S L7  
L16 16 S L8  
L17 372 S L9  
E NEUROPATHY  
L18 5490 S E3  
E EPILEPSY  
L19 10805 S E3-E12  
E SEIZURE  
L20 14562 S E3-E6  
E HYPOXIA  
L21 24801 S E3  
E STROKE  
L22 11880 S E3  
L23 0 S L17 AND L18  
L24 0 S L17 AND L19  
L25 0 S L17 AND L20  
L26 0 S L17 AND L21  
L27 0 S L17 AND L22  
L28 0 S L16 AND L18  
L29 25960 S L18 OR L19 OR L20  
L30 0 S L29 AND L16  
L31 0 S L29 AND L15  
L32 4 S L29 AND L14  
L33 0 S L29 AND L13  
L34 0 S L29 AND L12  
L35 30 S L29 AND L11  
L36 0 S L29 AND L10  
L37 350 S L21 AND L22  
L38 36331 S L21 OR L22  
L39 4 S L38 AND L10  
L40 2 S L38 AND L11  
L41 0 S L38 AND L12  
L42 1 S L38 AND L13  
L43 46 S L38 AND L14  
L44 1 S L38 AND L15  
L45 0 S L38 AND L16  
L46 0 S L38 AND L17  
E ISCHEMA  
L47 40127 S E6  
L48 3 S L47 AND L10  
L49 0 S L47 AND L16  
L50 3 S L47 AND L15  
L51 3 S L47 AND L11  
L52 0 S L47 AND L12  
L53 1 S L47 AND L13

L54 52 S L47 AND L14  
L55 28898 S HIS  
L56 47 S L54 NOT L43

=>

---Logging off of STN---

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Executing the logoff script...

=> LOG Y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	192.90	251.35
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-11.76	-11.76

STN INTERNATIONAL LOGOFF AT 16:08:56 ON 12 APR 2001